INTERIM DATA SUMMARY OF ONGOING BASELINE OFF-SITE AIR MONITORING VIA SAMPLING FOR VOLATILE ORGANIC COMPOUNDS AND HYDROGEN SULFIDE BY APPLICATION OF PASSIVE/DIFFUSIVE SAMPLING METHODS

WEST LAKE LANDFILL SITE BRIDGETON, MISSOURI CERCLIS ID: MOD079900932

Superfund Technical Assessment and Response Team (START) 4 Contract No. EP-S7-13-06, Task Order No. 0058



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EXECUTIVE SUMMARY

The Tetra Tech, Inc. (Tetra Tech) Superfund Technical Assessment and Response Team (START) has been tasked by the U.S. Environmental Protection Agency (EPA) to assist with baseline air monitoring at off-site locations around the West Lake Landfill site (WLLS) in Bridgeton, Missouri. This interim report summarizes results of sampling air for volatile organic compounds (VOC) and hydrogen sulfide (H₂S) by use of passive/diffusive samplers from December 18, 2014, through March 13, 2015. Baseline sampling via passive/diffusive methods was initiated during the baseline monitoring period after stakeholders had identified the sampling methodology for possible use during construction of the isolation barrier.

The baseline period air monitoring occurred at the following off-site monitoring stations according to the EPA-approved quality assurance project plan (QAPP):

Station 1 – Robertson Fire Protection District Station 2, 3820 Taussig Rd., Bridgeton, Missouri

Station 2 – Pattonville Fire Department District, 13900 St Charles Rock Rd., Bridgeton, Missouri

Station 3 – Pattonville Fire Department District Station 2, 3365 McKelvey Rd., Bridgeton, Missouri

Station 4 – Spanish Hillage Diagnostic Parish Spanish Parish Pa

Station 5 – St. Charles Fin Department Lation 2, 150 S. Jain St., St. Charles, Missouri.

The Station 1 through 4 locar ons were selected pronarily for the positions near and around WLLS (approximately 0.3 to 1 mile from WLLS, in various directions from WLLS). Station 5, designated as a reference (or background) station, is farther away from WLLS than the other stations, but still within the general vicinity so as to be representative of the North St. Louis County and eastern St. Charles County area.

Data from the passive/diffusive samplers collected from December 18, 2014, through March 13, 2015, were evaluated for differences in measured concentrations among the air monitoring stations off site of the WLLS. In addition, VOC results obtained by use of the passive/diffusive samplers were compared to VOC results previously obtained via 24-hour Summa® canister sampling at the WLLS air monitoring stations. The following describes the findings:

VOC concentrations detected at the five WLLS air monitoring stations via passive/diffusive sampling are within the concentration ranges previously detected at the air monitoring stations via 24-hour Summa canister sampling. Previous examination of 24-hour VOC concentrations detected at the five WLLS air monitoring stations revealed concentrations typical for an outdoor urban environment (see Tetra Tech 2015d). Thus, VOC concentrations detected at the five WLLS air monitoring stations via passive/diffusive sampling also appear typical for an outdoor urban environment.

- □ Statistical testing for differences in VOC concentrations detected via passive/diffusive sampling among the five air monitoring stations suggests tendency for detection of TCE at higher concentrations at Station 2 than at the other stations (Stations 1, 3, 4, and 5). A similar result was obtained from statistical testing of TCE data obtained via 24-hour sampling with Summa canisters at the five air monitoring stations. Although differences were detected among the stations, TCE concentrations at the five WLLS monitoring stations (including Station 2) were found comparable to 24-hour TCE concentrations EPA has detected at other urban areas via the National Air Toxics Treads Stations (NATTS) air monitoring program.
- No statistically significant differences in H_2S concentrations were found among the five WLLS air monitoring stations, and the concentrations were consistent with typical H_2S concentrations within outdoor urban environments.

Overall, the VOC and H₂S measurements obtained via passive/diffusive sampling at the off-site monitoring stations appear typical for outdoor urban measurements.

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DRAFT

1.0 INTRODUCTION

The Tetra Tech, Inc. (Tetra Tech) Superfund Technical Assessment and Response Team (START) has been tasked by the U.S. Environmental Protection Agency (EPA) to assist with baseline monitoring at off-site locations around the West Lake Landfill site (WLLS) in Bridgeton, Missouri. This interim report summarizes results from sampling air for volatile organic compounds (VOC) and hydrogen sulfide (H₂S) by use of passive/diffusive samplers from December 18, 2014, through March 13, 2015. Baseline sampling via passive/diffusive methods was initiated during the baseline monitoring period after stakeholders had identified the sampling methodology for possible use during construction of the isolation barrier.

For the overall baseline monitoring campaign, START's tasks have included: (1) assembling and maintaining a network of off-site air monitoring stations with instrumentation and sampling devices to measure parameters of concern, (2) collecting samples and coordinating laboratory analysis, (3) assisting EPA with data acquisition and management, (4) documenting off-site air monitoring efforts, and (5) validating/verifying initial screening of the data. The objective of this report is to present an interim summary of VOCs and massive acquired by use of practive/difference samplers, remaining findings related to data validation, validation and sability.

2.0 PROBLEM DEFINITION, BACKGROUND, AND SITE DESCRIPTION

EPA is conducting ongoing air monitoring at locations off site of WLLS during a pre-construction, baseline period prior to initiation of construction of a planned isolation barrier at WLLS. Air monitoring during the baseline period will provide data for use to (1) evaluate pre-construction concentrations of chemical and radiological parameters of potential concern in outdoor air, and (2) optimize the sampling and monitoring plan for the off-site air monitoring to occur during construction of the isolation barrier. During barrier construction, air monitoring will occur to address concerns that construction operations at WLLS could impact human health and the environment via release to ambient air of solid waste landfill gases of concern or of particulates with radiologically-impacted materials (RIM).

West Lake Landfill is an approximately 200-acre property including several closed solid waste landfill units that accepted wastes for landfilling from the 1940s or 1950s through 2004, plus a solid waste transfer station, a concrete plant, and an asphalt batch plant. WLLS is at 13570 St. Charles Rock Road in Bridgeton, St. Louis County, Missouri, approximately 1 mile north of the intersection of Interstate 70 and Interstate 270 (see Appendix A, Figure 1). WLLS was used for limestone quarrying and crushing begins ig in the operations from 193 te 1940 or carry 1950s, presons of the quarried ed or landf al refu industrial solutions, and areas and adjacent a as were ing m s of leached ba um sulfate residues (a construction/demoli on debri 973, a 0 to In oxin remnant from the Manuatan Engineer District Manuel Energy Commission project) was reportedly mixed with approximately 39,000 tons of soil from the 9200 Latty Avenue site in Hazelwood, Missouri, transported to the WLLS, and used as daily or intermediate cover material. In December 2004, the Bridgeton Sanitary Landfill—the last landfill unit to receive solid waste—stopped receiving waste pursuant to an agreement with the City of St. Louis to reduce potential for birds to interfere with Lambert Field International Airport operations. In December 2010, Bridgeton Landfill detected changes elevated temperatures and elevated carbon monoxide levels—in its landfill gas extraction system operating at the South Quarry of the Bridgeton Sanitary Landfill portion of the Site (a landfill portion not associated with known RIM). Further investigation indicated that the South Quarry Pit landfill was undergoing an exothermic subsurface smoldering event (SSE). In 2013, potentially responsible parties committed to constructing an isolation barrier that would separate the Bridgeton Landfill undergoing the SSE from the RIM-containing portions of WLLS (EPA 2014).

3.0 SAMPLING STRATEGY AND METHODOLOGY

In April/May 2014, EPA and START began baseline monitoring and sampling for the chemical and radiological parameters of concern at five off-site monitoring stations. Baseline sampling for VOCs and H₂S via passive/diffusive methods (the subject of this interim report) began in December 2014 after the sampling methodology had been identified for possible use during the isolation barrier construction. Sampling is occurring at the following monitoring stations (see Appendix A, Figure 1):

- Station 1 Robertson Fire Protection District Station 2, 3820 Taussig Rd., Bridgeton, Missouri
- Station 2 Pattonville Fire Department District, 13900 St Charles Rock Rd., Bridgeton, Missouri
- Station 3 Pattonville Fire Department District Station 2, 3365 McKelvey Rd., Bridgeton, Missouri
- Station 4 Spanish Village Park, 12827 Spanish Village Dr., Bridgeton, Missouri
- Station 5 St. Charles Fire Department Station #2, 1550 S. Main St., St. Charles, Missouri.

The Station 1 through 4 locations were selected primarily for their positions near and around WLLS (approximately 0.3) arious o ctions was designated as from. criterion that it be a reference (or back ound) st ion and its 1 lected according to th but still within the frequently upwind o WLLS a fa her a v fron e other station han orth St. Letis general vicinity so a ounty and east in St. Charles County area (see wind rose in Appendix A, Figure 1).

VOCs were identified as a parameter of potential concern in the Quality Assurance Project Plan (QAPP) (Tetra Tech 2014) based on historical information regarding the site and program experience with similar types of sites. Sampling for VOCs during the baseline monitoring period has occurred primarily by use of Summa® canisters (collected over a 24-hour period) and laboratory analysis via EPA Method TO-15. Sampling by use of Summa canisters occurred weekly at the air monitoring stations from May 8 to December 17, 2014, and the resulting data were summarized and evaluated by Tetra Tech (2015d). The report herein provides an interim summary of VOC sampling by use of passive/diffusive samplers that began in December 2014 after stakeholders had identified the sampling methodology for possible use during isolation barrier construction. This sampling occurred via deployment of Radiello® brand passive/diffusive samplers fitted with activated charcoal adsorbent cartridges. The cartridges were generally deployed at each of the five monitoring stations continuously for sampling durations of

approximately 7 days¹. At the end of the sampling duration, the deployed cartridges were collected, and a new sampling deployment was initiated with new cartridges. The Radiello cartridges for VOC sampling were shipped to Pace Analytical Laboratory (Pace) in Lenexa, Kansas, for analysis via Method EPA TO-17 modified for analysis of the Radiello cartridges.

H₂S, another parameter of potential concern identified in the QAPP (Tetra Tech 2014), was also sampled by use of Radiello cartridges concurrently with the VOC Radiello sampling at the five air monitoring stations. Radiello samplers deployed for H₂S sampling were fitted with adsorbent cartridges containing zinc acetate. These cartridges were shipped to ALS Laboratory in Simi Valley, California, for analysis via an extraction and colorimetric analysis specified by the Radiello cartridge manufacturer.



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¹ The Radiello cartridges collected on January 19, 2015 were deployed for 14 days to assess for any significant differences in measured analyte concentrations (or detection limits) compared to a 7 day deployment (no significant differences were observed).

4.0 SUMMARY AND EVALUATION OF VOC AND H₂S RESULTS

The following sections present data summaries of the VOC and H₂S results from Radiello passive/diffusive samplers during the baseline monitoring period, including time series and box plots of the data, and results of statistical analyses.

4.1 DATA VALIDATION, VERIFICATION, AND USABILITY

As laboratory analytical reports are received for the analyses, START reviews and qualifies the data according to the EPA Contract Laboratory Program National Functional Guidelines for Superfund Organic Methods Data Review guidance document (EPA 2008). This is standard practice for EPA data. A data validation report is appended to each analytical laboratory report, and is included in each data deliverable package (see Tetra Tech 2015a, b, c, d). Qualifications to the data from START's review are indicated by qualifier flags that accompany the data presented therein. Overall, review of the laboratory analytical data packages indicated that quality of the VOC and H₂S data was acceptable and usable as qualified for the intended purposes of the data.

On March 19, 2015. tra Tec mists h identified ben ene and toluene ed hat its contamination in the orbent n Irbent media supplied by the ared to te fro ne or toluene oncentrations. The manufacturer. For t sample affe ted, Pa ben ot repor affected samples were generally those collected on and after January 28, 2015. Because the potentially affected data (benzene and toluene concentrations) were not reported by the laboratory, no specific qualifications were assigned via START's data validation/verification.

4.2 RESULTS AND EVALUATION

The following describes results of sampling for VOCs and H₂S from December 18, 2014, through March 13, 2015, by use of passive/diffusive Radiello samplers.

4.2.1 Summary of Results

Appendix B presents the Radiello sampling results and summary statistics for VOCs (see Tables B-1 through B-14) and H₂S (see Table B-15). Appendix C presents a time-series plot for each VOC (see Exhibits C-1 through C-14) and for H₂S (see Exhibit C-15). As discussed in Section 4.2.3, the time-series

plots for the VOCs also show, for comparison, VOC results obtained via 24-hour sampling by use of Summa canisters².

4.2.2 Comparison of Results Among Off-Site Monitoring Stations

Differences in measured concentrations of VOCs and H₂S among the air monitoring stations are evaluated and described in this section so that data users can be aware of those differences when using the preconstruction baseline data. Comparisons of measured concentrations among the air monitoring stations proceeded via statistical testing and examination of boxplots. Results obtained by use of Radiello cartridges were compared (comparisons of Summa VOC results among the stations appears in Tetra Tech 2015e). The following describes the evaluations.

Kruskal-Wallis Statistical Test

The Kruskal-Wallis statistical test was used to test for differences in VOCs and H₂S concentrations among the five monitoring stations. The Kruskal-Wallis test compares multiple treatments (such as the multiple monitoring locations), and was selected in particular because it is a non-parametric statistical (rank- or percentiles surts found in the data. Before perfor ance of t te the dat ed for ie Kruskal-Wa s test as recommended in Statistics for Cen. red Env ntal I d R (Helsel 20 were censored at the report g limit it hadataset by assuming all values below the highest reporting limit (including all non-detects and any reported value less than the highest reporting limit) a low and arbitrary value; the value "-1" was used. The statistical software package R was used to conduct the Kruskal-Wallis tests. Results of the analysis suggested statistically significant differences in concentrations of one analyte among the five monitoring stations:

Trichloroethene (TCE): Measurements of concentration at Station 2 tended to be higher than at Stations 1, 3, 4, and 5.

The analysis found no statistically significant difference in H₂S concentrations among the five monitoring stations. A summary of the Kruskal-Wallis test results is in Appendix D, Table D-1.

The same statistical analysis applied to VOC results obtained via 24-hour sampling by use of Summa canisters also found statistically significant differences in TCE concentrations among the five stations (the

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² Data Summary of Ongoing Baseline Off-Site Air Monitoring, Volatile Organic Compounds by Test Method TO-15 (Tetra Tech 2015d) presents an evaluation of VOC results obtained via 24-hour sampling by use of Summa canisters.

test indicated that measurements of concentration at Station 2 tended to be higher than at Stations 1, 3, and 4) (see Tetra Tech 2015e).

Boxplots

Boxplots of VOC and H₂S results were constructed and evaluated to verify results of the Kruskal-Wallis statistical testing (see Appendix E). Boxplots are visual comparisons of data that display relative positions of the 25th, 50th, and 75th percentiles, and also individual outlier data points. The "NADA" (Nondetects and Data Analysis for environmental data) for the statistical software package R was used to create "censored" boxplots of the VOC data. In constructing the censored boxplots, the NADA software accounts for non-detect values and displays a horizontal line across the boxplots representing the maximum "less than" value in the data. Boxplot elements above the line are statistically accurate, but boxplot elements below the line represent only estimated percentiles (based on the distribution of the uncensored data). Review of the boxplots supports findings of the Kruskal-Wallis testing that indicated no significant statistical differences in measured concentrations of H₂S or VOCs (except for TCE) among the five monitoring stations.

4.2.3 Compariso of VOC Results from adiell and Sum a Canister Methods

To compare passive liffusive ladia to sandling deal with late 10m previous 24 nour sampling by use of Summa canisters³, but that sets were plotted at the Appendix Coplots (isopropy enzene, methyl-tert-butyl ether, and *trans*-1,2-dichloroethene were reported only in the Radiello analysis). Notably, for those VOCs concentrations measured by both methods, Radiello results were within the range of concentrations measured by use of Summa canisters.

4.2.4 Comparison of H₂S Results to Typical Ambient Concentrations

The reported H_2S concentrations ranged 0.16 to 0.64 micrograms per cubic meter ($\mu g/m^3$) (approximately 0.12 to 0.46 parts per billion [ppb]). The Agency for Toxic Substances and Disease Registry (ATSDR) reports that ambient air concentrations of H_2S from natural sources range between 0.11 and 0.33 ppb, and that concentrations of H_2S in urban areas are generally less than 1 ppb (ATSDR 2014). Thus, H_2S results from the five off-site monitoring stations appear typical for outdoor urban measurements.

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³ Data Summary of Ongoing Baseline Off-Site Air Monitoring, Volatile Organic Compounds by Test Method TO-15 (Tetra Tech 2015d) presents an evaluation of VOC results obtained via 24-hour sampling by use of Summa canisters.

5.0 ANALYSIS OF RESULTS

As discussed above, statistical testing for differences in concentrations among the five air monitoring stations suggests a tendency for measurements of TCE concentration to be higher at Station 2 than at the other stations 1, 3, 4, and 5. A similar result was obtained from statistical testing of the TCE data obtained via 24-hour sampling by use of Summa canisters at the five air monitoring stations; however, the 24-hour TCE concentrations were found comparable to 24-hour TCE concentrations EPA had detected at other urban areas under the National Air Toxics Treads Stations (NATTS) air monitoring program (see Tetra Tech 2015e). TCE concentrations detected via Radiello sampling are within the range of 24-hour concentrations previously detected at WLLS air monitoring stations via 24-hour Summa sampling (see Appendix C, Exhibit C-13), and are thus also comparable to concentrations detected in other urban areas.

TCE is a common industrial solvent that frequently appears in groundwater contaminant plumes at cleanup sites from historical use and improper disposal. Given the chemical properties of TCE, if found in the buried waste at WLLS, TCE likely would be found in the groundwater and released to the air. However, EPA reviewed groundwater data from WLLS and found no indication of presence of TCE. would be detected in the leachate Further, if TCE wer at WL it like the t rial However ΓCE was also not resulting from mois re percolating brough e buri te ma d at V LS pr lec . Thus, TCE des not appear to be detected in leachate imples of tme at WLLS, and, by inference, the letections in ambient air are not believed to present in the buried have resulted from releases of this VOC from WLLS. Moreover, EPA's examination of reports of air emissions from various industrial facilities in the area revealed presence of other possible sources of TCE.

6.0 SUMMARY OF OBSERVATIONS

VOCs and H₂S sampling via 1- and 2-week deployments of passive/diffusive samplers began in December 2014 as part of baseline off-site air monitoring at five air monitoring stations off site of WLLS. This interim report summarizes and evaluates data from the passive/diffusive samplers obtained from December 18, 2014, through March 13, 2015. Differences in measured concentrations among the air monitoring stations off site of the WLLS were evaluated. In addition, VOC results obtained by use of the passive/diffusive samplers were compared to VOC results previously obtained via 24-hour Summa canister sampling at the WLLS air monitoring stations. The following describes the findings.

- VOC concentrations detected at the five WLLS air monitoring stations via passive/diffusive sampling are within the concentration ranges previously detected at the air monitoring stations via 24-hour Summa canister sampling. Previous examination of 24-hour VOC concentrations detected at the five WLLS air monitoring stations revealed concentrations typical for an outdoor urban environment (see Tetra Tech 2015e). Thus, VOC concentrations detected at the five WLLS air monitoring stations via passive/diffusive sampling also appear typical for an outdoor urban environment.
- Statistical testing for differences in VOC concentrations detected via passive/diffusive sampling ror detection of real at higher among the f nitor endenc sugges). A similar result was concentrations at Stati s (Stations 1, 3, 4, and n 2 an at th other i obtained from statistic tes E dat obta ed vi r sample g with Summa erences were th d tected among the Almo canisters at e five ai toring tions moi stations, TC ration at the fi LLS monoring stations (including Station 2) were found comparable to 24-hour TCE concentrations EPA had detected at other urban areas via the National Air Toxics Treads Stations (NATTS) air monitoring program.
- No statistically significant differences in H_2S concentrations were found among the five WLLS air monitoring stations, and the concentrations were consistent with typical H_2S concentrations within outdoor urban environments.

Overall, the VOCs and H₂S measurements obtained via passive/diffusive sampling at the off-site monitoring stations appear typical for outdoor urban measurements.

7.0 REFERENCES

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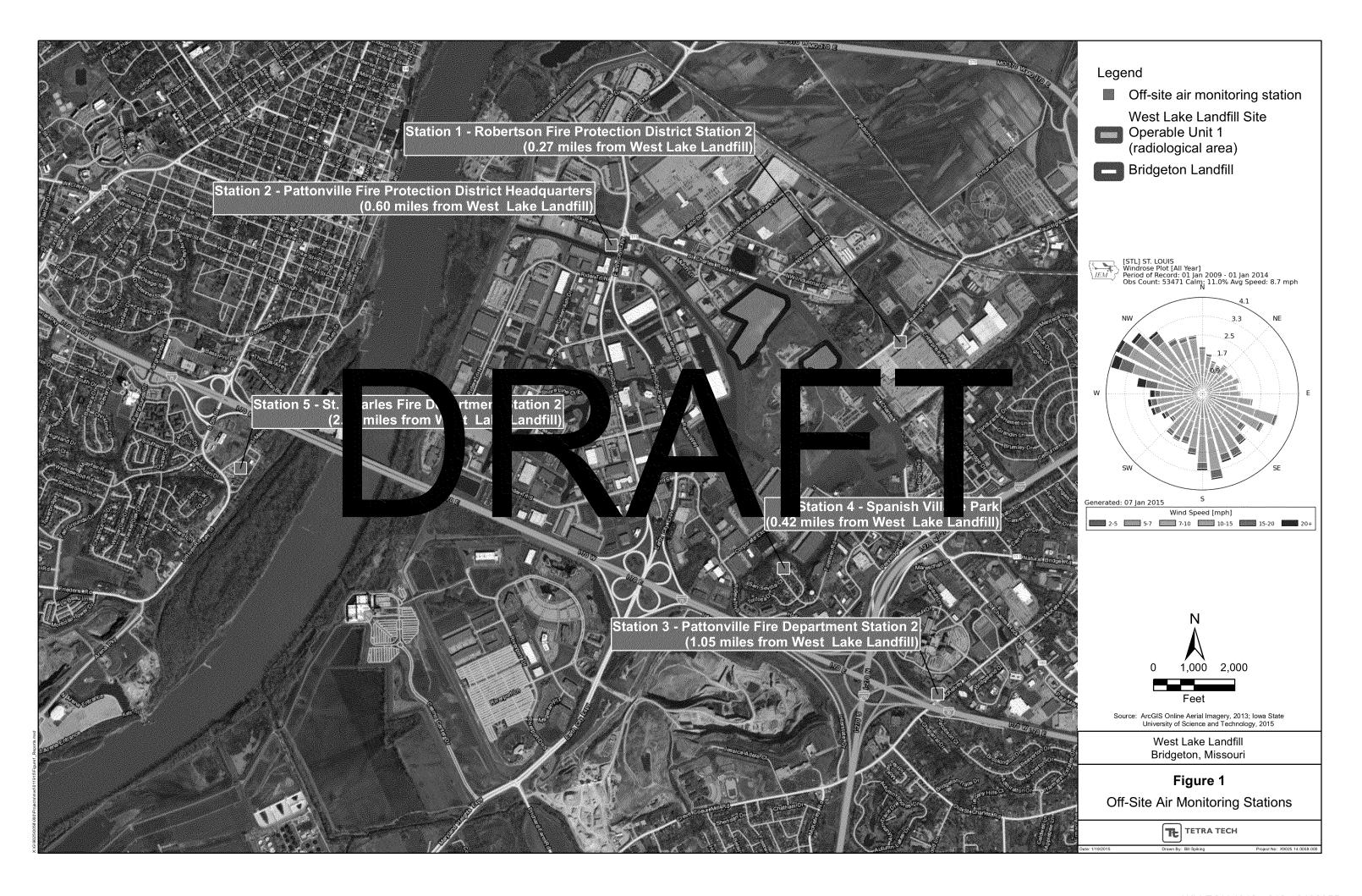






Table B-1
1,2,4-Trimethylbenzene

Date Collected	Station 1	Station 2	Station 3	Station 4	Station 5
12/18/2014	ND (0.012)	ND (0.013)	ND (0.013)	ND (0.012)/ND (0.012)	ND (0.013)
12/27/2014	ND (0.012)	ND (0.012)	ND (0.012)	ND (0.012)/ND (0.012)	ND (0.012)
1/5/2015	ND (0.011)	ND (0.011)	ND (0.011)	ND (0.011)/ND (0.011)	ND (0.011)
1/19/2015	ND (0.0074)	ND (0.0074)	ND (0.0074)	ND (0.0074)/ND (0.0074)	ND (0.0074)
1/28/2015	0.26	0.27	0.32	0.25 /0.35	0.45
2/4/2015	ND (0.034)	ND (0.034)	ND (0.034)	ND (0.034)/ND (0.034)	ND (0.034)
2/13/2015	ND (0.031)/ND (0.026)	ND (0.026)	ND (0.026)	ND (0.026)	ND (0.026)
2/20/2015	U	U	U	U/U	U
2/27/2015	ND (0.14)	ND (0.14)	ND (0.14)	ND (0.14)/ND (0.14)	ND (0.14)
3/6/2015	ND (0.14)	ND (0.14)	ND (0.14)	ND (0.14)/ND (0.14)	ND (0.14)
3/13/2015	ND (0.15)	ND (0.15)	ND (0.15)	ND (0.14)/ND (0.14)	ND (0.15)
No. of Detects	1	1	1	2	1
No. of Samples	12	11	11	21	11
Minimum	0.26	0.27	0.32	0.25	0.45
Median	0.26	0.27	0.32	0.3	0.45
Maximum	0.26	0.27	0.32	0.35	0.45

All concentrations in micrograms per cubic meter $(\mu g/m^{3})$ ND () not detected (method detection limit)



Table B-2 1,3,5-Trimethylbenzene

Date Collected	Station 1	Station 2	Station 3	Station 4	Station 5
12/18/2014	ND (0.012)	ND (0.012)	ND (0.012)	ND (0.012)/ND (0.012)	ND (0.012)
12/27/2014	ND (0.011)	ND (0.011)	ND (0.011)	ND (0.011)/ND (0.011)	ND (0.011)
1/5/2015	ND (0.01)	ND (0.011)	ND (0.011)	ND (0.011)/ND (0.011)	ND (0.011)
1/19/2015	ND (0.0068)	ND (0.0068)	ND (0.0068)	ND (0.0068)/ND (0.0068)	ND (0.0068)
1/28/2015	ND (0.032)	ND (0.032)	ND (0.032)	ND (0.032)/ND (0.032)	ND (0.032)
2/4/2015	ND (0.041)	ND (0.041)	ND (0.041)	ND (0.041)/ND (0.041)	ND (0.041)
2/13/2015	ND (0.038)/ND (0.032)	ND (0.032)	ND (0.032)	ND (0.032)	ND (0.032)
2/20/2015	ND (0.041)	ND (0.041)	U	ND (0.041)/U	ND (0.041)
2/27/2015	ND (0.17)	ND (0.17)	ND (0.17)	ND (0.17)/ND (0.17)	ND (0.17)
3/6/2015	ND (0.17)	ND (0.17)	ND (0.17)	ND (0.17)/ND (0.17)	ND (0.17)
3/13/2015	ND (0.18)	ND (0.18)	ND (0.18)	ND (0.17)/ND (0.17)	ND (0.18)
No. of Detects	0	0	0	0	0
No. of Samples	12	11	11	21	11
Minimum	NA	NA	NA	NA	NA
Median	NA	NA	NA	NA	NA
Maximum	NA	NA	NA	NA	NA

All concentrations in micrograms per cubic meter (µg/m³)

NA not applicable

ND () not detected (method detection limit)



Table B-3 Benzene

Date Collected	Station 1	Station 2	Station 3	Station 4	Station 5
12/18/2014	ND (0.039)	ND (0.039)	ND (0.039)	ND (0.039)/ND (0.039)	ND (0.039)
12/27/2014	ND (0.036)	ND (0.036)	ND (0.036)	ND (0.036)/ND (0.036)	ND (0.036)
1/5/2015	ND (0.035)	ND (0.035)	ND (0.035)	ND (0.035)/ND (0.035)	ND (0.035)
1/19/2015	ND (0.023)	ND (0.023)	ND (0.023)	ND (0.023)/ND (0.023)	ND (0.023)
2/20/2015	NR	NR	U	NR/U	NR
No. of Detects	0	0	0	0	0
No. of Samples	4	4	5	9	4
Minimum	NA	NA	NA	NA	NA
Median	NA	NA	NA	NA	NA
Maximum	NA	NA	NA	NA	NA

All concentrations in micrograms per cubic meter ($\mu g/m^3$)

NA not applicable

ND () not detected (method detection limit)

NR not reported by laboratory



Table B-4 cis-1,2-Dichloroethene

Date Collected	Station 1	Station 2	Station 3	Station 4	Station 5
12/18/2014	ND (0.076)	ND (0.076)	ND (0.076)	ND (0.076)/ND (0.076)	ND (0.076)
12/27/2014	ND (0.071)	ND (0.071)	ND (0.071)	ND (0.071)/ND (0.071)	ND (0.071)
1/5/2015	ND (0.068)	ND (0.069)	ND (0.069)	ND (0.069)/ND (0.069)	ND (0.069)
1/19/2015	ND (0.045)	ND (0.045)	ND (0.045)	ND (0.045)/ND (0.045)	ND (0.045)
1/28/2015	ND (0.025)	ND (0.025)	ND (0.025)	ND (0.025)/ND (0.025)	ND (0.025)
2/4/2015	ND (0.032)	ND (0.032)	ND (0.032)	ND (0.032)/ND (0.032)	ND (0.032)
2/13/2015	ND (0.029)/ND (0.025)	ND (0.025)	ND (0.025)	ND (0.025)	ND (0.025)
2/20/2015	ND (0.032)	ND (0.032)	U	ND (0.032)/U	ND (0.032)
2/27/2015	ND (0.13)	ND (0.13)	ND (0.13)	ND (0.13)/ND (0.13)	ND (0.13)
3/6/2015	ND (0.13)	ND (0.13)	ND (0.13)	ND (0.13)/ND (0.13)	ND (0.13)
3/13/2015	ND (0.14)	ND (0.14)	ND (0.14)	ND (0.14)/ND (0.14)	ND (0.14)
No. of Detects	0	0	0	0	0
No. of Samples	12	11	11	21	11
Minimum	NA	NA	NA	NA	NA
Median	NA	NA	NA	NA	NA
Maximum	NA	NA	NA	NA	NA

All concentrations in micrograms per cubic meter (µg/m³)

NA not applicable

ND () not detected (method detection limit)



Table B-5 Ethylbenzene

Date Collected	Station 1	Station 2	Station 3	Station 4	Station 5
12/18/2014	ND (0.01)	ND (0.01)	ND (0.01)	ND (0.01)/ND (0.01)	ND (0.01)
12/27/2014	ND (0.0096)	ND (0.0096)	ND (0.0096)	ND (0.0096)/ND (0.0096)	ND (0.0096)
1/5/2015	ND (0.0092)	ND (0.0093)	ND (0.0093)	ND (0.0092)/ND (0.0092)	ND (0.0093)
1/19/2015	ND (0.006)	ND (0.006)	ND (0.006)	ND (0.006)/ND (0.006)	ND (0.006)
1/28/2015	0.32	0.31	0.29	0.28 /0.31	0.37
2/4/2015	0.19	0.18	0.18	0.18 /0.16	0.17
2/13/2015	0.21 /0.19	0.19	0.2	0.19	0.26
2/20/2015	U	U	U	U/U	U
2/27/2015	ND (0.08)	ND (0.08)	ND (0.08)	ND (0.08)/ND (0.08)	ND (0.08)
3/6/2015	ND (0.08)	ND (0.08)	ND (0.08)	ND (0.08)/ND (0.08)	ND (0.08)
3/13/2015	ND (0.082)	ND (0.082)	ND (0.082)	ND (0.081)/ND (0.081)	ND (0.082)
No. of Detects	4	3	3	5	3
No. of Samples	12	11	11	21	11
Minimum	0.19	0.18	0.18	0.16	0.17
Median	0.2	0.19	0.2	0.19	0.26
Maximum	0.32	0.31	0.29	0.31	0.37

All concentrations in micrograms per cubic meter $(\mu g/m^{3})$ ND () not detected (method detection limit)



Table B-6 Isopropylbenzene (Cumene)

Date Collected	Station 1	Station 2	Station 3	Station 4	Station 5
12/18/2014	ND (0.011)	ND (0.011)	ND (0.011)	ND (0.011)/ND (0.011)	ND (0.011)
12/27/2014	ND (0.011)	ND (0.011)	ND (0.011)	ND (0.011)/ND (0.011)	ND (0.011)
1/5/2015	ND (0.01)	ND (0.01)	ND (0.01)	ND (0.01)/ND (0.01)	ND (0.01)
1/19/2015	ND (0.0066)	ND (0.0066)	ND (0.0066)	ND (0.0066)/ND (0.0066)	ND (0.0066)
1/28/2015	ND (0.018)	ND (0.018)	ND (0.018)	ND (0.018)/ND (0.018)	ND (0.018)
2/4/2015	ND (0.023)	ND (0.023)	ND (0.023)	ND (0.023)/ND (0.023)	ND (0.023)
2/13/2015	ND (0.021)/ND (0.018)	ND (0.018)	ND (0.018)	ND (0.018)	ND (0.018)
2/20/2015	ND (0.023)	ND (0.023)	U	ND (0.023)/U	ND (0.023)
2/27/2015	ND (0.096)	ND (0.096)	ND (0.096)	ND (0.096)/ND (0.096)	ND (0.096)
3/6/2015	ND (0.096)	ND (0.096)	ND (0.096)	ND (0.096)/ND (0.096)	ND (0.096)
3/13/2015	ND (0.099)	ND (0.099)	ND (0.099)	ND (0.098)/ND (0.098)	ND (0.099)
No. of Detects	0	0	0	0	0
No. of Samples	12	11	11	21	11
Minimum	NA	NA	NA	NA	NA
Median	NA	NA	NA	NA	NA
Maximum	NA	NA	NA	NA	NA

Notes:

All concentrations in micrograms per cubic meter (µg/m³)

NA not applicable

ND () not detected (method detection limit)



Table B-7 m&p-Xylene

Date Collected	Station 1	Station 2	Station 3	Station 4	Station 5
12/18/2014	ND (0.025)	ND (0.025)	ND (0.025)	ND (0.025)/ND (0.025)	ND (0.025)
12/27/2014	ND (0.023)	ND (0.023)	ND (0.023)	ND (0.023)/ND (0.023)	ND (0.023)
1/5/2015	ND (0.022)	ND (0.023)	ND (0.023)	ND (0.023)/ND (0.023)	ND (0.023)
1/19/2015	ND (0.015)	ND (0.015)	ND (0.015)	ND (0.015)/ND (0.015)	ND (0.015)
1/28/2015	0.98	0.96	0.89	0.84 /0.91	1.1
2/4/2015	0.5	0.48	0.47	0.47 /0.41	0.44
2/13/2015	0.49 J/0.51 J	0.49 J	0.52 J	0.47 J	0.66 J
2/20/2015	U	U	U	U/U	U
2/27/2015	ND (0.07)	ND (0.07)	ND (0.07)	ND (0.07)/ND (0.07)	ND (0.07)
3/6/2015	ND (0.07)	ND (0.07)	ND (0.07)	ND (0.07)/ND (0.07)	ND (0.07)
3/13/2015	ND (0.071)	ND (0.071)	ND (0.071)	0.87 /0.94	0.87
No. of Detects	4	3	3	7	4
No. of Samples	12	11	11	21	11
Minimum	0.49	0.48	0.47	0.41	0.44
Median	0.505	0.49	0.52	0.84	0.765
Maximum	0.98	0.96	0.89	0.94	1.1

All concentrations in micrograms per cubic meter ($\mu g/m^3$)

J laboratory flag indicating an estimated result less than reporting limit

ND () not detected (method detection limit)



Table B-8 Methyl-tert-butyl ether

Date Collected	Station 1	Station 2	Station 3	Station 4	Station 5
12/18/2014	ND (0.033)	ND (0.033)	ND (0.033)	ND (0.033)/ND (0.033)	ND (0.033)
12/27/2014	ND (0.031)	ND (0.031)	ND (0.031)	ND (0.031)/ND (0.031)	ND (0.031)
1/5/2015	ND (0.03)	ND (0.03)	ND (0.03)	ND (0.03)/ND (0.03)	ND (0.03)
1/19/2015	ND (0.02)	ND (0.02)	ND (0.02)	ND (0.02)/ND (0.02)	ND (0.02)
1/28/2015	ND (0.035)	ND (0.035)	ND (0.035)	ND (0.035)/ND (0.035)	ND (0.035)
2/4/2015	ND (0.045)	ND (0.045)	ND (0.045)	ND (0.045)/ND (0.045)	ND (0.045)
2/13/2015	ND (0.042)/ND (0.035)	ND (0.035)	ND (0.035)	ND (0.035)	ND (0.035)
2/20/2015	ND (0.046)	ND (0.046)	U	ND (0.046)/U	ND (0.046)
2/27/2015	ND (0.19)	ND (0.19)	ND (0.19)	ND (0.19)/ND (0.19)	ND (0.19)
3/6/2015	ND (0.19)	ND (0.19)	ND (0.19)	ND (0.19)/ND (0.19)	ND (0.19)
3/13/2015	ND (0.2)	ND (0.2)	ND (0.2)	ND (0.2)/ND (0.2)	ND (0.2)
No. of Detects	0	0	0	0	0
No. of Samples	12	11	11	21	11
Minimum	NA	NA	NA	NA	NA
Median	NA	NA	NA	NA	NA
Maximum	NA	NA	NA	NA	NA

All concentrations in micrograms per cubic meter ($\mu g/m^3$)

J laboratory flag indicating an estimated result less than reporting limit

NA not applicable

ND () not detected (method detection limit)



Table B-9 o-Xylene

Date Collected	Station 1	Station 2	Station 3	Station 4	Station 5
12/18/2014	ND (0.014)	ND (0.014)	ND (0.014)	ND (0.014)/ND (0.014)	ND (0.014)
12/27/2014	ND (0.014)	ND (0.014)	ND (0.014)	ND (0.014)/ND (0.014)	ND (0.014)
1/5/2015	ND (0.013)	ND (0.013)	ND (0.013)	ND (0.013)/ND (0.013)	ND (0.013)
1/19/2015	ND (0.0085)	ND (0.0085)	ND (0.0085)	ND (0.0085)/ND (0.0085)	ND (0.0085)
1/28/2015	0.34	0.33	0.3	0.29 /0.31	0.39
2/4/2015	0.18	0.17	0.17	0.17 /0.15	0.16
2/13/2015	0.18 /0.18	0.18	0.19	0.17	0.24
2/20/2015	U	U	U	U/U	U
2/27/2015	ND (0.083)	ND (0.083)	ND (0.083)	ND (0.083)/ND (0.083)	ND (0.083)
3/6/2015	ND (0.083)	ND (0.083)	ND (0.083)	ND (0.083)/ND (0.083)	ND (0.083)
3/13/2015	ND (0.085)	ND (0.085)	ND (0.085)	ND (0.085)/ND (0.085)	ND (0.085)
No. of Detects	4	3	3	5	3
No. of Samples	12	11	11	21	11
Minimum	0.18	0.17	0.17	0.15	0.16
Median	0.18	0.18	0.19	0.17	0.24
Maximum	0.34	0.33	0.3	0.31	0.39

All concentrations in micrograms per cubic meter $(\mu g/m^{3})$ ND () not detected (method detection limit)



Table B-10 Tetrachloroethene

Date Collected	Station 1	Station 2	Station 3	Station 4	Station 5
12/18/2014	ND (0.032)	ND (0.032)	ND (0.032)	ND (0.031)/ND (0.031)	ND (0.032)
12/27/2014	ND (0.03)	ND (0.03)	ND (0.03)	ND (0.03)/ND (0.03)	ND (0.03)
1/5/2015	ND (0.029)	ND (0.029)	ND (0.029)	ND (0.029)/ND (0.029)	ND (0.029)
1/19/2015	ND (0.019)	ND (0.019)	ND (0.019)	ND (0.019)/ND (0.019)	ND (0.019)
1/28/2015	0.27	0.46	ND (0.03)	ND (0.03)/ND (0.03)	0.23
2/4/2015	ND (0.039)	ND (0.039)	ND (0.039)	ND (0.039)/ND (0.039)	ND (0.039)
2/13/2015	ND (0.036)/ND (0.03)	ND (0.03)	ND (0.03)	ND (0.03)	ND (0.03)
2/20/2015	ND (0.039)	ND (0.039)	U	ND (0.039)/U	ND (0.039)
2/27/2015	ND (0.17)	ND (0.17)	ND (0.17)	ND (0.17)/ND (0.17)	ND (0.17)
3/6/2015	ND (0.17)	ND (0.16)	ND (0.17)	ND (0.17)/ND (0.17)	ND (0.17)
3/13/2015	ND (0.17)	ND (0.17)	ND (0.17)	ND (0.17)/ND (0.17)	ND (0.17)
No. of Detects	1	1	0	0	1
No. of Samples	12	11	11	21	11
Minimum	0.27	0.46	NA	NA	0.23
Median	0.27	0.46	NA	NA	0.23
Maximum	0.27	0.46	NA	NA	0.23

All concentrations in micrograms per cubic meter (µg/m³)

NA not applicable

ND () not detected (method detection limit)



Table B-11 Toluene

Date Collected	Station 1	Station 2	Station 3	Station 4	Station 5
12/18/2014	ND (0.017)	ND (0.017)	ND (0.017)	ND (0.017)/ND (0.017)	ND (0.017)
12/27/2014	ND (0.016)	ND (0.016)	ND (0.016)	ND (0.016)/ND (0.016)	ND (0.016)
1/5/2015	ND (0.016)	ND (0.016)	ND (0.016)	ND (0.016)/ND (0.016)	ND (0.016)
1/19/2015	0.58	0.56	ND (0.01)	0.56 J/0.50	0.39 J
2/20/2015	NR	NR	0.24 J, CL, L2	0.20 J, CL, L2	NR
No. of Detects	1	1	1	3	1
No. of Samples	4	4	5	9	4
Minimum	0.58	0.56	0.24	0.2	0.39
Median	0.58	0.56	0.24	0.5	0.39
Maximum	0.58	0.56	0.24	0.56	0.39

All concentrations in micrograms per cubic meter (µg/m³)

CL indicates the continue calibration for the compound was outside of the laboratory's acceptance limits and the result may be biased low L2 indicates analyte recovery in the laboratory control sample was below quality control limits and the result may be biased low

ND () not detected (method detection limit)

NR not reported by laboratory



Table B-12 trans-1,2-Dichloroethene

Date Collected	Station 1	Station 2	Station 3	Station 4	Station 5
12/18/2014	ND (0.033)	ND (0.033)	ND (0.033)	ND (0.033)/ND (0.033)	ND (0.033)
12/27/2014	ND (0.031)	ND (0.031)	ND (0.031)	ND (0.031)/ND (0.031)	ND (0.031)
1/5/2015	ND (0.03)	ND (0.03)	ND (0.03)	ND (0.03)/ND (0.03)	ND (0.03)
1/19/2015	ND (0.019)	ND (0.019)	ND (0.019)	ND (0.019)/ND (0.019)	ND (0.019)
1/28/2015	ND (0.019)	ND (0.019)	ND (0.019)	ND (0.019)/ND (0.019)	ND (0.019)
2/4/2015	ND (0.024)	ND (0.024)	ND (0.024)	ND (0.024)/ND (0.024)	ND (0.024)
2/13/2015	ND (0.022)/ND (0.019)	ND (0.019)	ND (0.019)	ND (0.019)	ND (0.019)
2/20/2015	ND (0.024)	ND (0.024)	U	ND (0.024)/U	ND (0.024)
2/27/2015	ND (0.1)	ND (0.1)	ND (0.1)	ND (0.1)/ND (0.1)	ND (0.1)
3/6/2015	ND (0.1)	ND (0.1)	ND (0.1)	ND (0.1)/ND (0.1)	ND (0.1)
3/13/2015	ND (0.1)	ND (0.1)	ND (0.1)	ND (0.1)/ND (0.1)	ND (0.1)
No. of Detects	0	0	0	0	0
No. of Samples	12	11	11	21	11
Minimum	NA	NA	NA	NA	NA
Median	NA	NA	NA	NA	NA
Maximum	NA	NA	NA	NA	NA

All concentrations in micrograms per cubic meter (µg/m³)

NA not applicable

ND () not detected (method detection limit)



Table B-13 Trichloroethene

Date Collected	Station 1	Station 2	Station 3	Station 4	Station 5
12/18/2014	ND (0.031)	ND (0.031)	ND (0.031)	ND (0.031)/ND (0.031)	ND (0.031)
12/27/2014	ND (0.029)	ND (0.029)	ND (0.029)	ND (0.029)/ND (0.029)	ND (0.029)
1/5/2015	ND (0.028)	ND (0.028)	ND (0.028)	ND (0.028)/ND (0.028)	ND (0.028)
1/19/2015	ND (0.018)	ND (0.018)	ND (0.018)	ND (0.018)/ND (0.018)	ND (0.018)
1/28/2015	ND (0.026)	0.51	ND (0.026)	ND (0.026)/ND (0.026)	ND (0.026)
2/4/2015	ND (0.033)	ND (0.033)	ND (0.033)	ND (0.033)/ND (0.033)	ND (0.033)
2/13/2015	ND (0.031)/ND (0.026)	0.33	ND (0.026)	ND (0.026)	ND (0.026)
2/20/2015	ND (0.033)	0.41	U	ND (0.033)/U	ND (0.033)
2/27/2015	ND (0.14)	ND (0.14)	ND (0.14)	ND (0.14)/ND (0.14)	ND (0.14)
3/6/2015	ND (0.14)	ND (0.14)	ND (0.14)	ND (0.14)/ND (0.14)	ND (0.14)
3/13/2015	ND (0.14)	ND (0.14)	ND (0.14)	ND (0.14)/ND (0.14)	ND (0.14)
No. of Detects	0	3	0	0	0
No. of Samples	12	11	11	21	11
Minimum	NA	0.33	NA	NA	NA
Median	NA	0.41	NA	NA	NA
Maximum	NA	0.51	NA	NA	NA

All concentrations in micrograms per cubic meter (µg/m³)

NA not applicable

ND () not detected (method detection limit)



Table B-14 Vinyl chloride

Date Collected	Station 1	Station 2	Station 3	Station 4	Station 5
12/18/2014	ND (0.22)	ND (0.22)	ND (0.22)	ND (0.22)/ND (0.22)	ND (0.22)
12/27/2014	ND (0.2)	ND (0.2)	ND (0.2)	ND (0.2)/ND (0.2)	ND (0.2)
1/5/2015	ND (0.19)	ND (0.2)	ND (0.2)	ND (0.2)/ND (0.2)	ND (0.2)
1/19/2015	ND (0.13)	ND (0.13)	ND (0.13)	ND (0.13)/ND (0.13)	ND (0.13)
1/28/2015	ND (0.014)	ND (0.014)	ND (0.014)	ND (0.014)/ND (0.014)	ND (0.014)
2/4/2015	ND (0.019)	ND (0.019)	ND (0.019)	ND (0.019)/ND (0.019)	ND (0.019)
2/13/2015	ND (0.017)/ND (0.014)	ND (0.014)	ND (0.014)	ND (0.014)	ND (0.014)
2/20/2015	ND (0.019)	ND (0.019)	U	ND (0.019)/U	ND (0.019)
2/27/2015	ND (0.078)	ND (0.078)	ND (0.078)	ND (0.078)/ND (0.078)	ND (0.078)
3/6/2015	ND (0.078)	ND (0.078)	ND (0.078)	ND (0.078)/ND (0.078)	ND (0.078)
3/13/2015	ND (0.08)	ND (0.08)	ND (0.08)	ND (0.08)/ND (0.08)	ND (0.08)
No. of Detects	0	0	0	0	0
No. of Samples	12	11	11	21	11
Minimum	NA	NA	NA	NA	NA
Median	NA	NA	NA	NA	NA
Maximum	NA	NA	NA	NA	NA

All concentrations in micrograms per cubic meter $(\mu g/m^3)$

NA not applicable

ND () not detected (method detection limit)



Table B-15 Hydrogen Sulfide

Date Collected	Station 1	Station 2	Station 3	Station 4	Station 5
12/18/2014	0.57 Ј	0.56 J	0.54 J	0.57 J/0.51 J	0.64 J
12/27/2014	0.40 Ј	0.49 J	0.46 J	0.40 J/0.59 J	0.51 J
1/5/2015	0.53 J	0.37 J	0.55 J	0.60 J/0.62 J	0.55 J
1/19/2015	0.29 Ј	0.22 J	0.16 J	0.30 J/0.24 J	0.24 J
1/28/2015	0.45 J	0.39 J	0.38 J	0.59 J/0.41 J	0.25 J
2/4/2015	0.31 J	0.24 J	ND (0.21)	0.28 J/0.28 J	0.26 J
2/13/2015	ND (0.17)/ND (0.17)	ND (0.17)	ND (0.17)	ND (0.17)	ND (0.17)
2/20/2015	ND (0.25)	0.39 J	ND (0.25)	ND (0.25)/ND (0.25)	ND (0.25)
2/27/2015	ND (0.24)	ND (0.24)	ND (0.24)	ND (0.24)/ND (0.24)	ND (0.24)
3/6/2015	ND (0.23)	ND (0.23)	ND (0.23)	ND (0.23)/0.24 J	ND (0.23)
3/13/2015	0.24 J	0.48 J	ND (0.2)	0.43 J/0.38 J	0.31 J
No. of Detects	7	8	5	15	7
No. of Samples	12	11	11	21	11
Minimum	0.24	0.22	0.16	0.24	0.24
Median	0.4	0.39	0.46	0.41	0.31
Maximum	0.57	0.56	0.55	0.62	0.64

All concentrations in micrograms per cubic meter ($\mu g/m^3$)

J laboratory flag indicating an estimated result less than reporting limit

ND () not detected (method detection limit)





Exhibit C - 1 1,2,4 - Trimethylbenzene

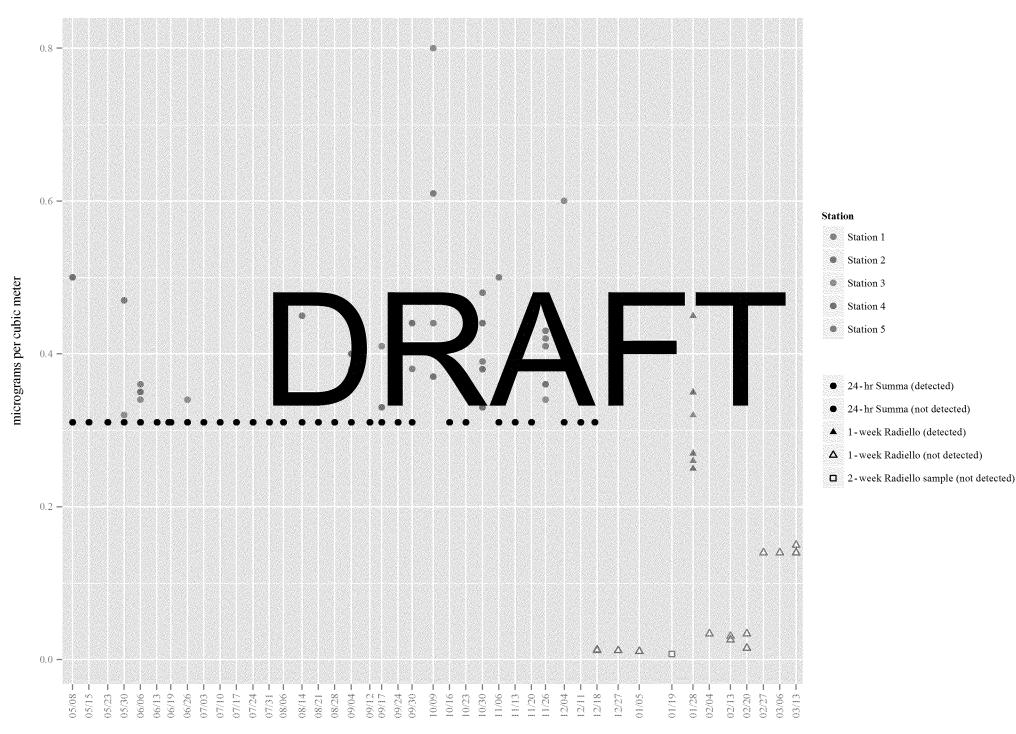


Exhibit C-2 1,3,5-Trimethylbenzene

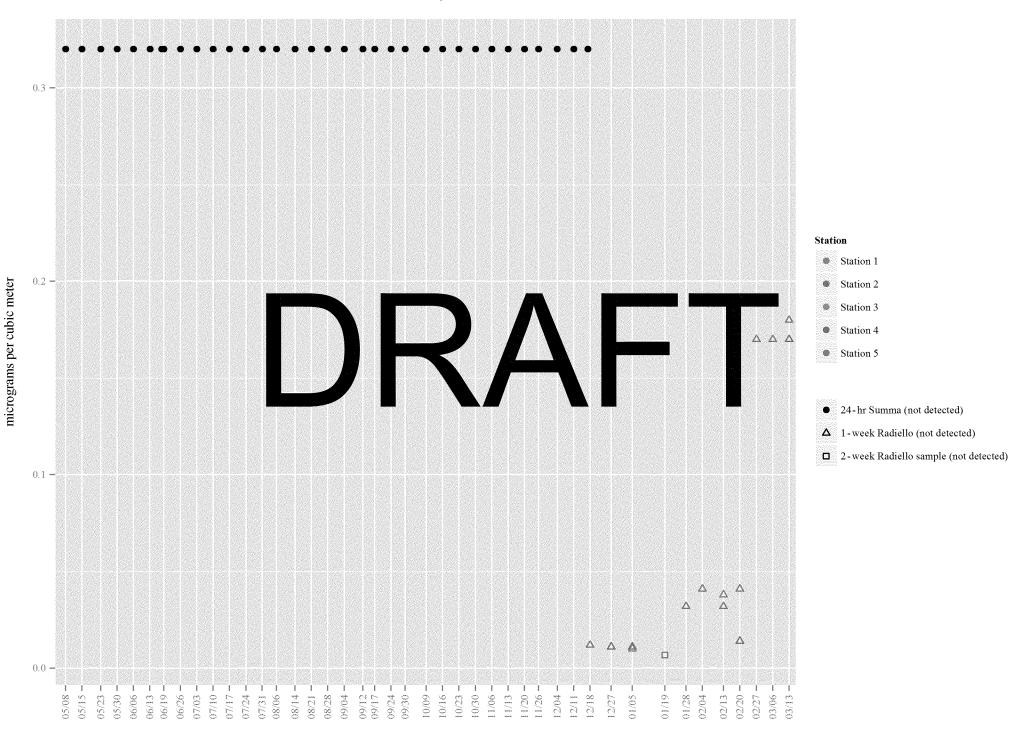


Exhibit C-3
Benzene

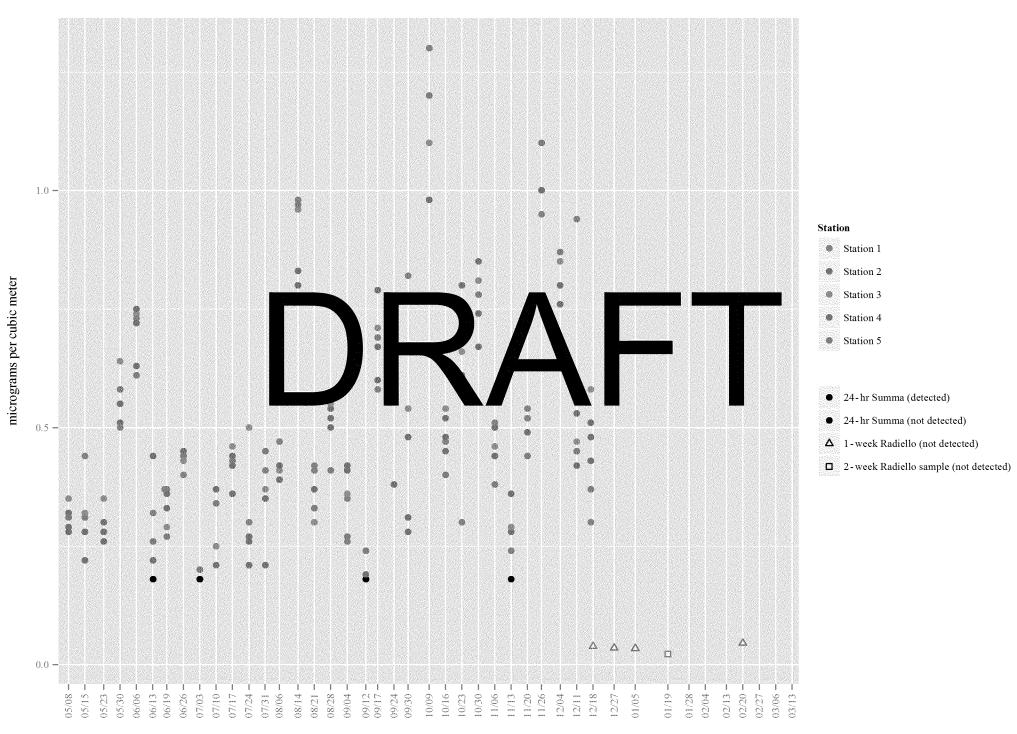


Exhibit C-4 cis-1,2-Dichloroethene

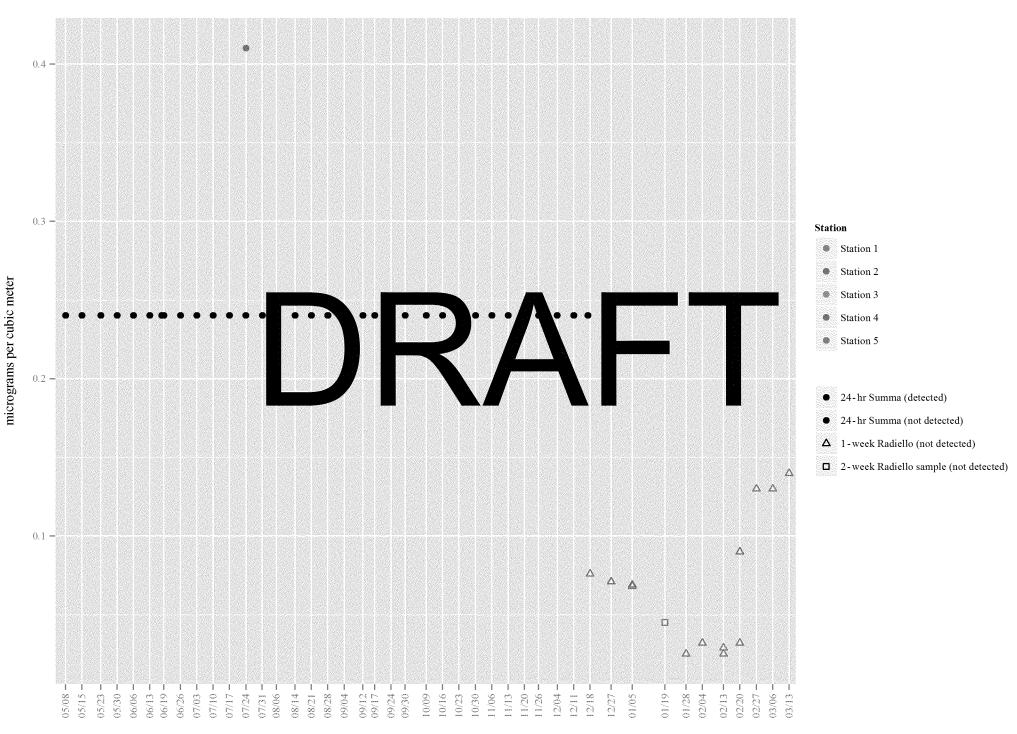


Exhibit C-5 Ethylbenzene

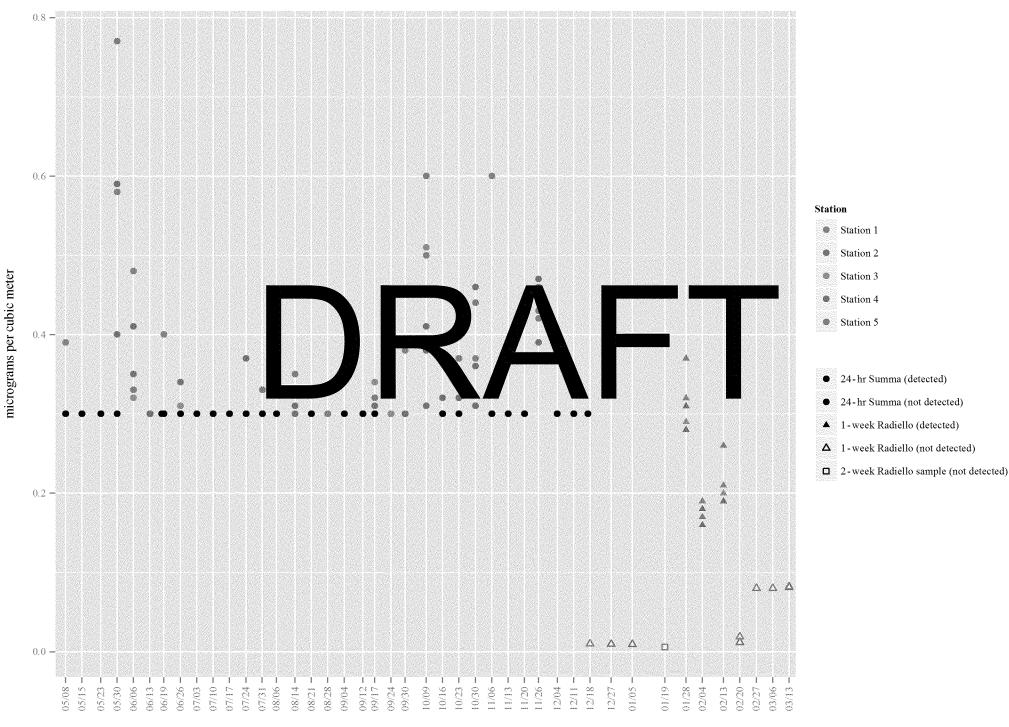


Exhibit C-6
Isopropylbenzene (Cumene)

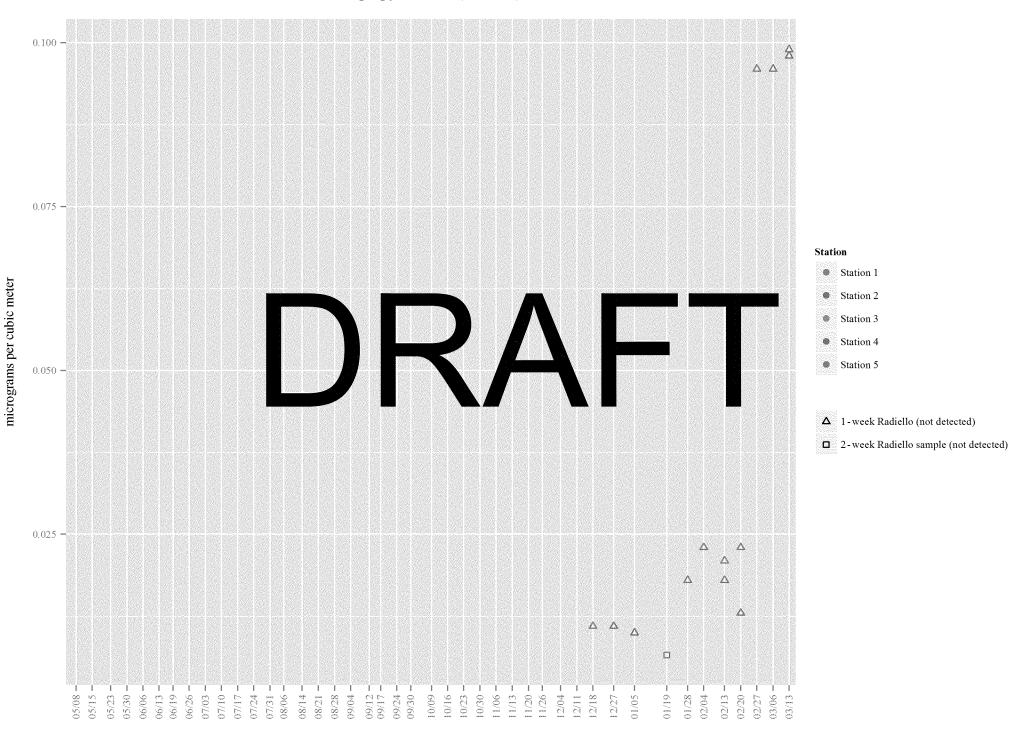


Exhibit C-7 m&p-Xylene

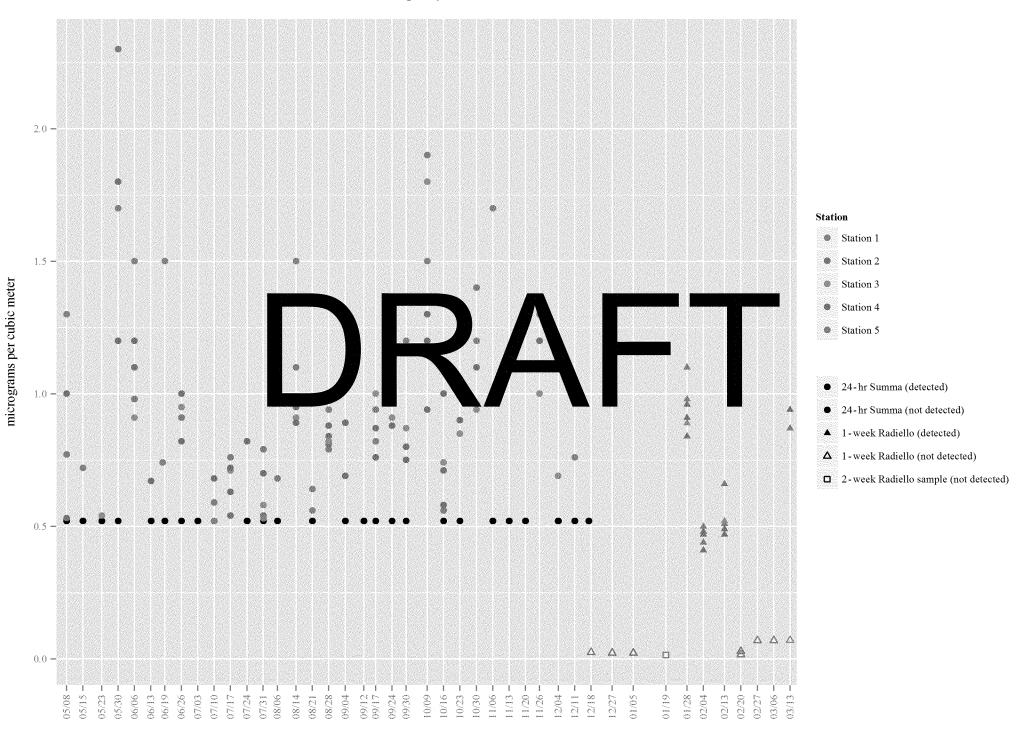


Exhibit C-8 Methyl-tert-butyl ether

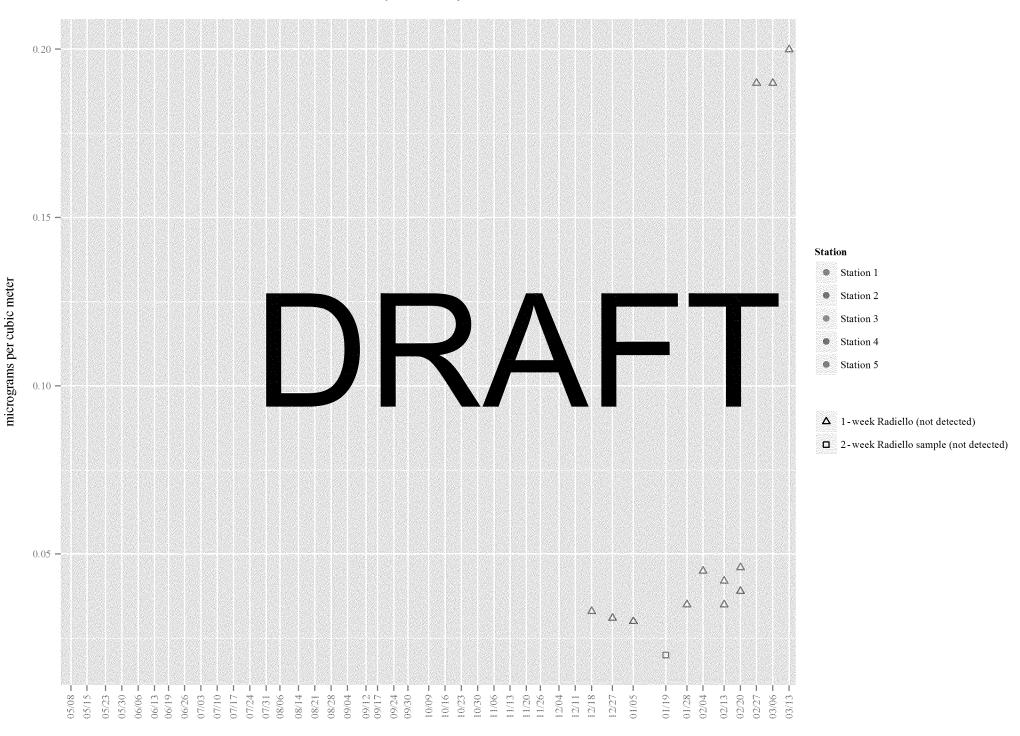


Exhibit C-9 o-Xylene

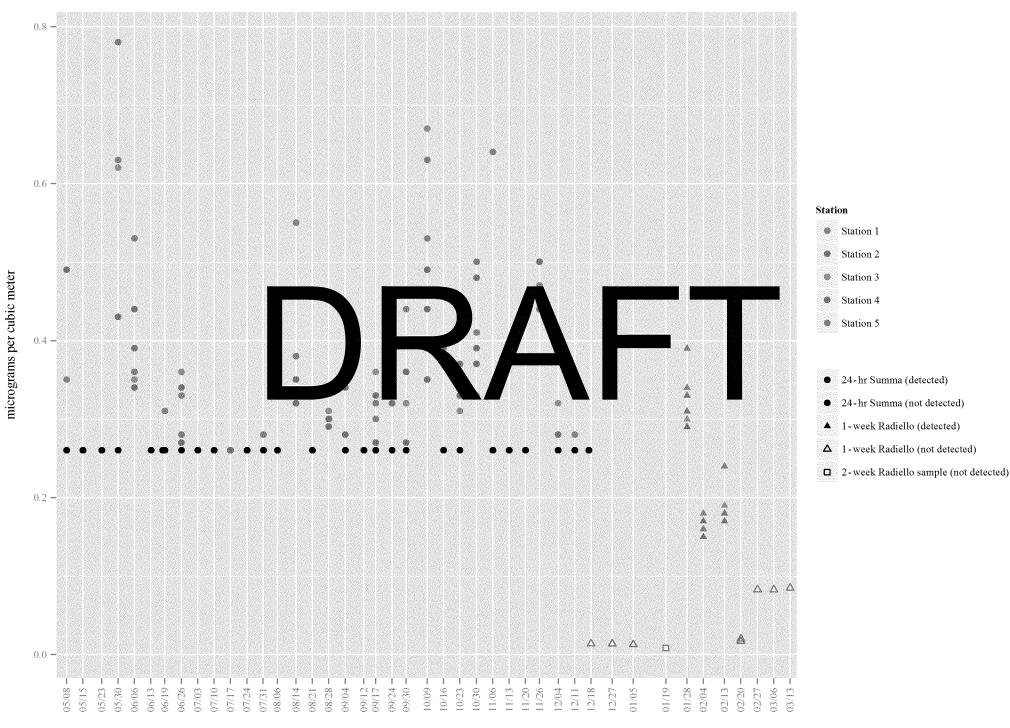


Exhibit C-10 Tetrachloroethene

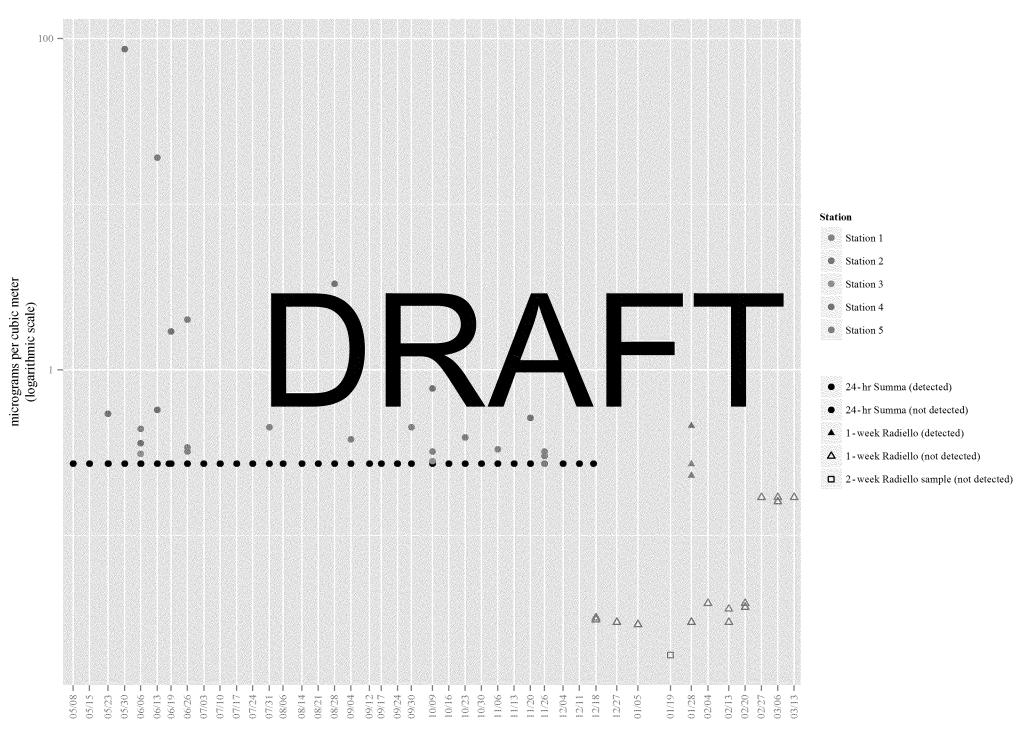
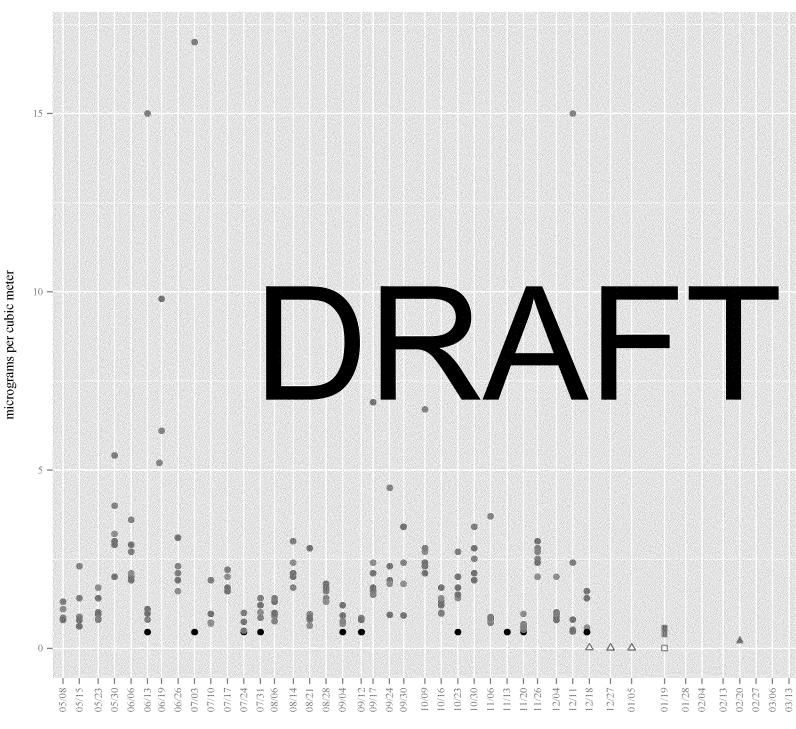


Exhibit C-11 Toluene



Station

- Station 1
- Station 2
- Station 3
- Station 4
- Station 5
- 24-hr Summa (detected)
- 24-hr Summa (not detected)
- ▲ 1-week Radiello (detected)
- Δ 1-week Radiello (not detected)
- 2-week Radiello (detected)
- ☐ 2-week Radiello sample (not detected)

Date (2014/2015)

Exhibit C - 12 trans - 1,2 - Dichloroethene

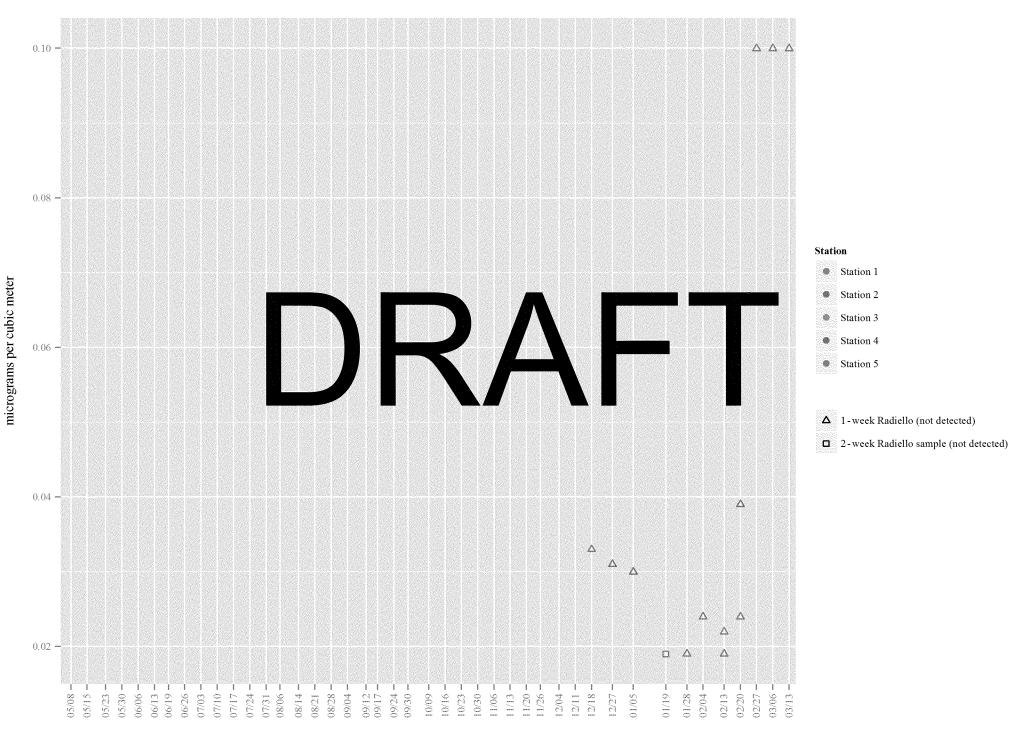


Exhibit C-13 Trichloroethene

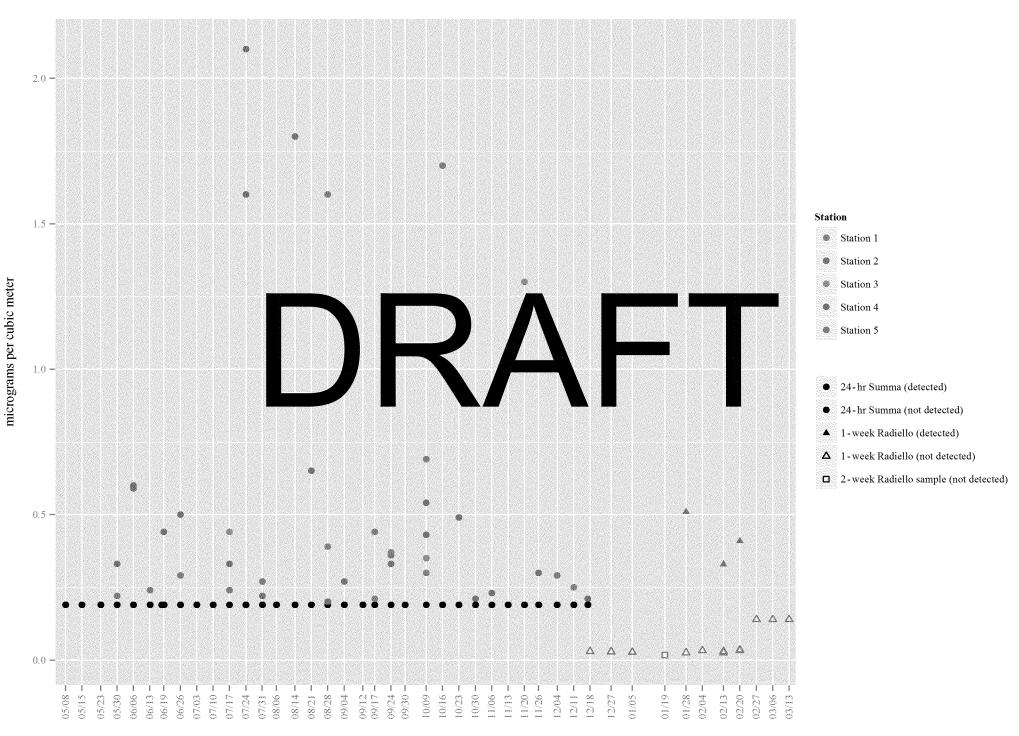
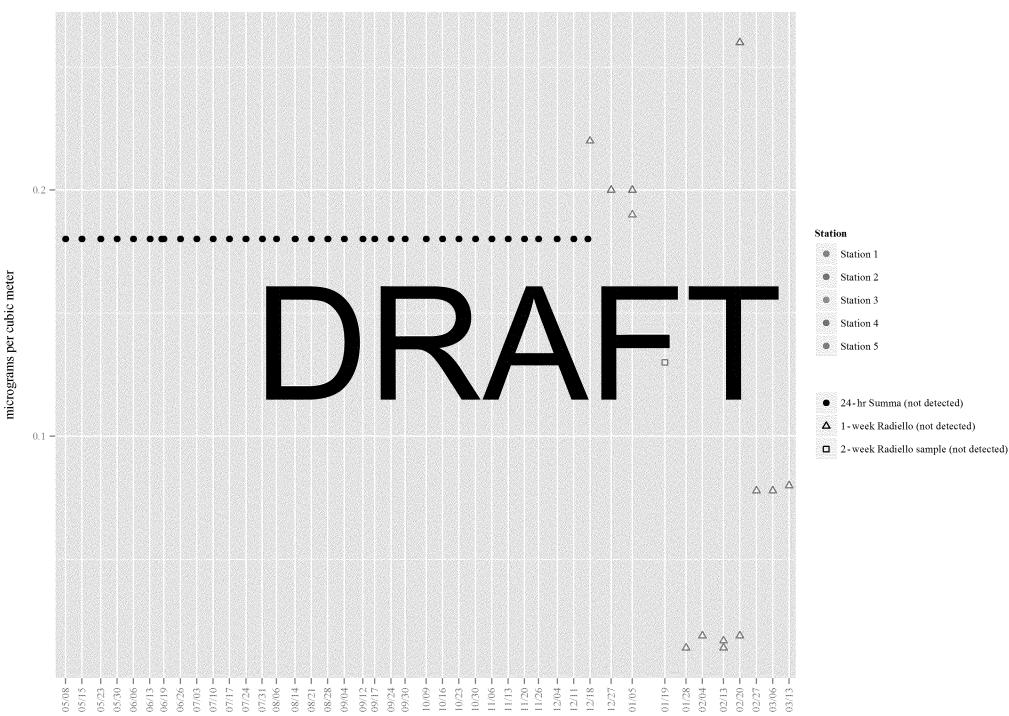
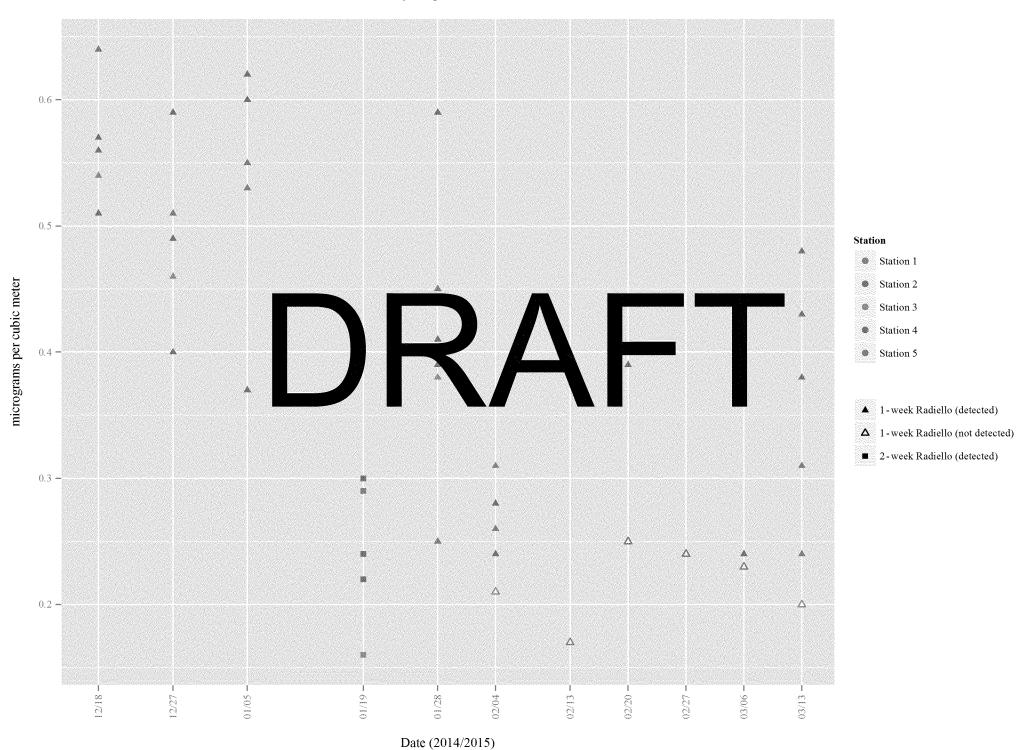


Exhibit C-14 Vinyl chloride



- △ 1-week Radiello (not detected)

Exhibit C-15 Hydrogen Sulfide



PPER DIX D FREQUENCY OF DETECTION SEMMAN STATISTICAL ANALYSIS RESULTS

TABLE D-1 COMPARISON OF PASSIVE/DIFFUSIVE SAMPLING RESULTS AT OFF-SITE AIR MONITORING STATIONS WEST LAKE LANDFILL, BRIDGETON, MISSOURI

Analyte ¹	Frequency of Detection ²						Results of Statistical Comparison Among
	Station 1	Station 2	Station 3	Station 4	Station 5	Total	Off-Site WLLS Stations ^{3, 4}
Hydrogen Sulfide	58.3%	72.7%	45.5%	71.4%	63.6%	62.3%	No significant difference $(p\text{-}value = 0.7549)$
m&p-Xylene	33.3%	27.3%	27.3%	33.3%	36.4%	31.5%	No significant difference $(p\text{-}value = 0.9777)$
Ethylbenzene	33.3%	27.3%	27.3%	23.8%	27.3%	27.8%	No significant difference $(p\text{-value} = 0.9722)$
o-Xylene	33.3%	27.3%	27.3%	23.8%	27.3%	27.8%	No significant difference $(p\text{-}value = 0.9688)$
Toluene	25.0%	25.0%	20.0%	33.3%	25.0%	25.7%	No significant difference $(p\text{-}value = 0.9819)$
1,2,4-Trimethylbenzene	8.3%	9.1%	9.1%	9.5%	9.1%	9.0%	No significant difference $(p\text{-}value = 0.9999)$
Trichloroethene	0.0%	27.3%	0.0%	0.0%	0.0%	5.5%	Station 2 tended to have higher measurements than the other stations $(p-value = 1.858e-05)$
Tetrachloroethene	8.3%	9.1%	0.0%	0.0%	9.1%	5.3%	No significant difference (p-value = 0.6012)
1,3,5-Trimethylbenzene	0.0%	0.0%	0.0%	0.0%	0.0%	0.0%	Analyte not detected
Benzene	0.0%	0.0%	0.0%	0.0%	0.0%	0.0%	Analyte not detected
cis -1,2-Dichloroethene	0.0%	0.0%	0.0%	0.0%	0.0%	0.0%	Analyte not detected
Isopropylbenzene (Cumene)	0.0%	0.0%	0.0%	0.0%	0.0%	0.0%	Analyte not detected
Methyl-tert-butyl ether	0.00/	0.0%	0.00/	0.0%	0.0%	0.0%	Analyte not detected
trans -1,2-Dichloroethene	0%	0.	(%	0.0	0.0%	0%	Analyt ot detected
Vinyl chloride	0%	0.0	(%	0.09	0.0%	%	Analyte ot detected

Notes

Analytes listed in descending ask according to overall percent sect of

² Frequency of detection is the rate of analyte detection via laboratory analysis of the passive/diffusive samplers. Detection of an analyte in either the sample or its duplicate was counted as one detect.

³ Results from the statistical software package R version 3.1.2 using the non-parametric Kruskal-Wallis test to compare mean/median characteristics of compounds among monitoring stations. A p-value equal to or less than 0.05 suggests significant differences in mean/median characteristics among the stations. A p-value exceeding 0.05 suggests that mean/median characteristics among stations are comparable.

⁴ Shading indicates detection of a difference.



BOXPLOT DESCRIPTION AND KEY

Boxplots render visual comparisons of data by displaying relative positions of the 25th, 50th, and 75th percentiles, and also individual outlier data points. The "NADA" (Nondetects and Data Analysis for environmental data) for the statistical software package R was used to create "censored" boxplots of the volatile organic compound (VOC) data. In constructing the censored boxplots, the NADA software accounts for non-detect values and displays a horizontal line across the boxplots representing the maximum "less than" value in the data. Boxplot elements above the line are statistically accurate, but boxplot elements below the line represent only estimated percentiles (based on the distribution of the uncensored data).

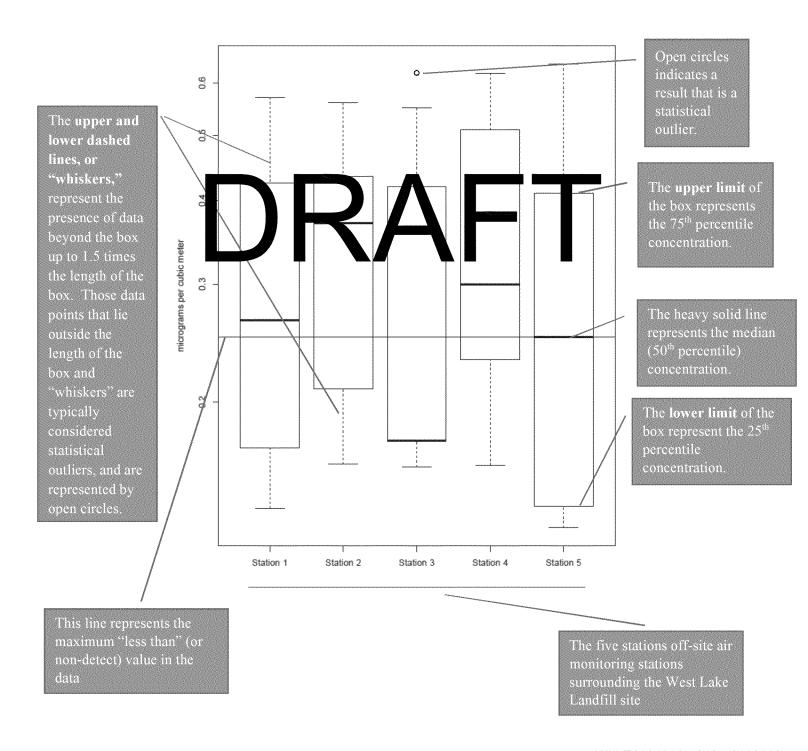


Exhibit E-1 1,2,4-Trimethylbenzene

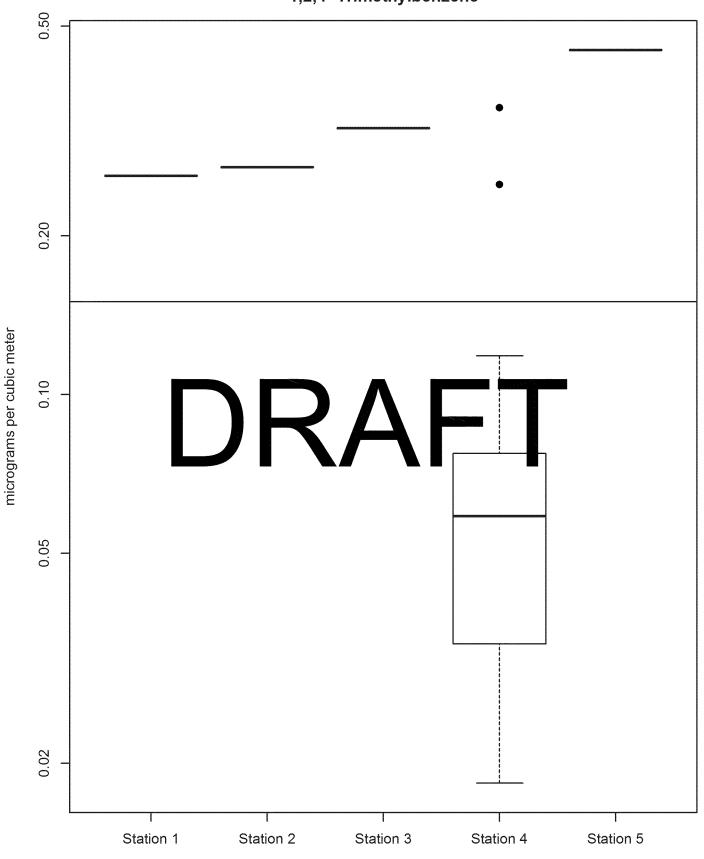


Exhibit E-2 Ethylbenzene

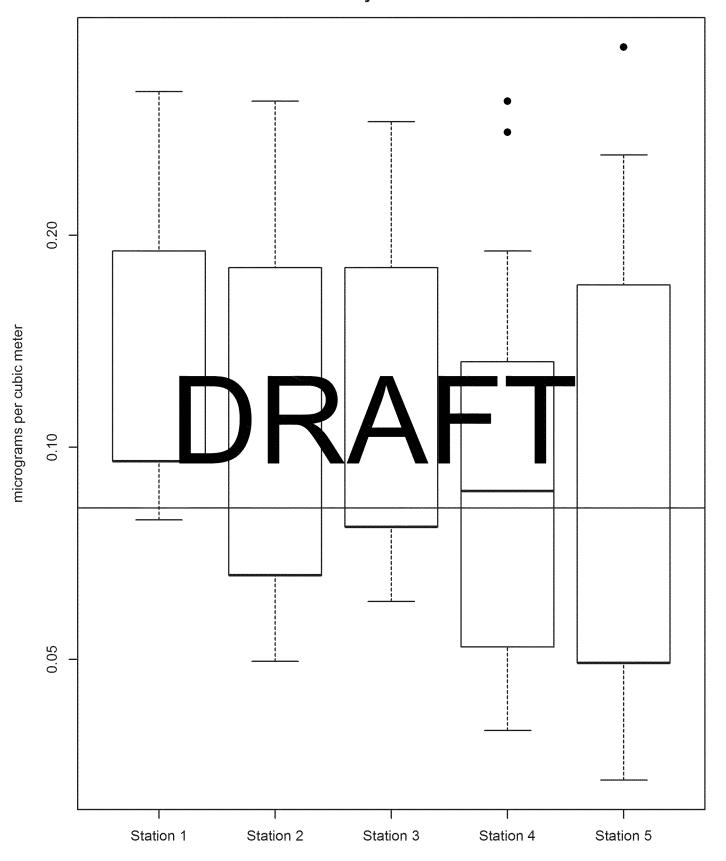


Exhibit E-3 Hydrogen Sulfide

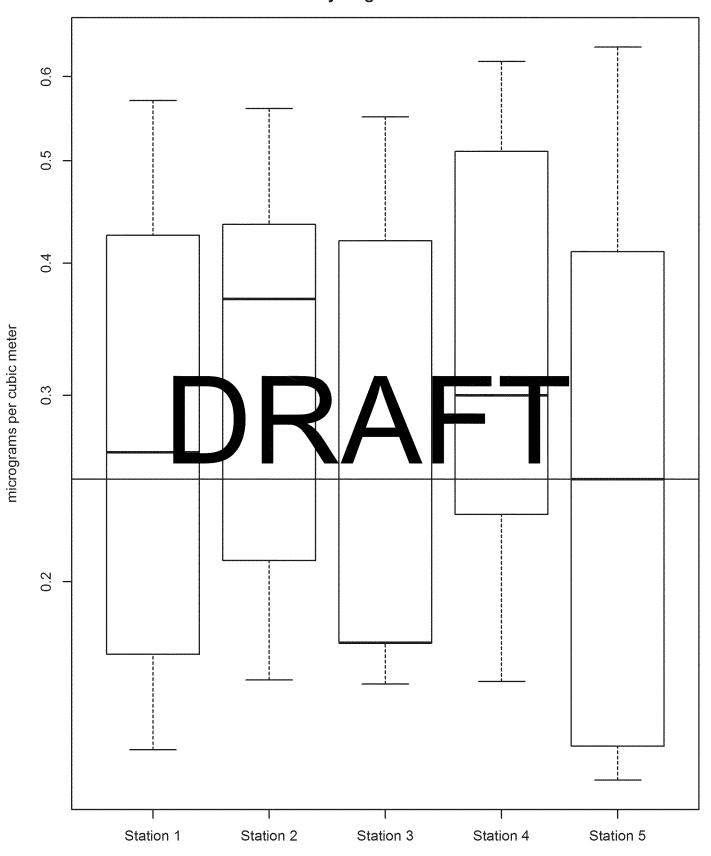


Exhibit E-4 m&p-Xylene

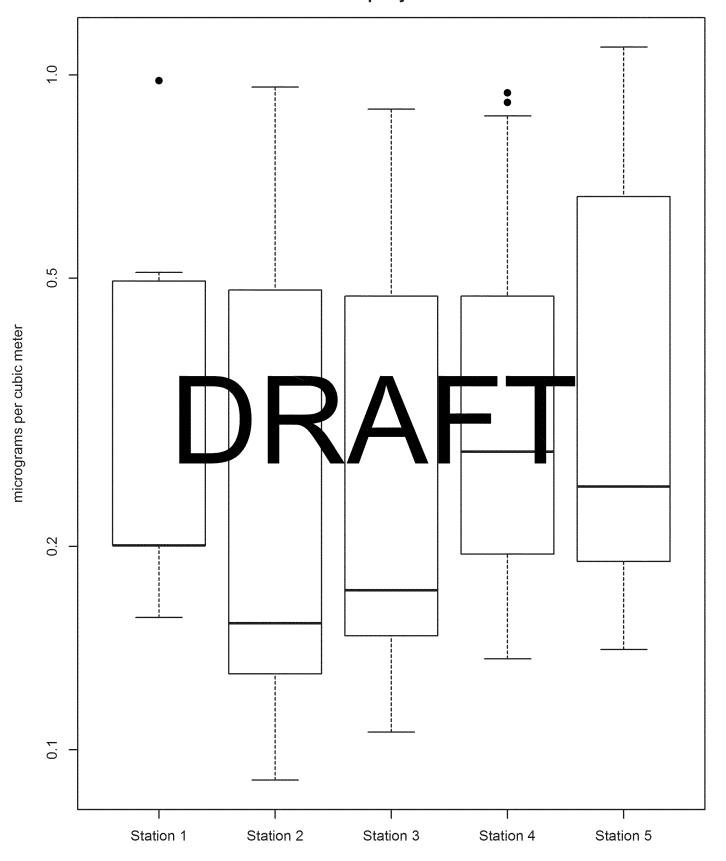
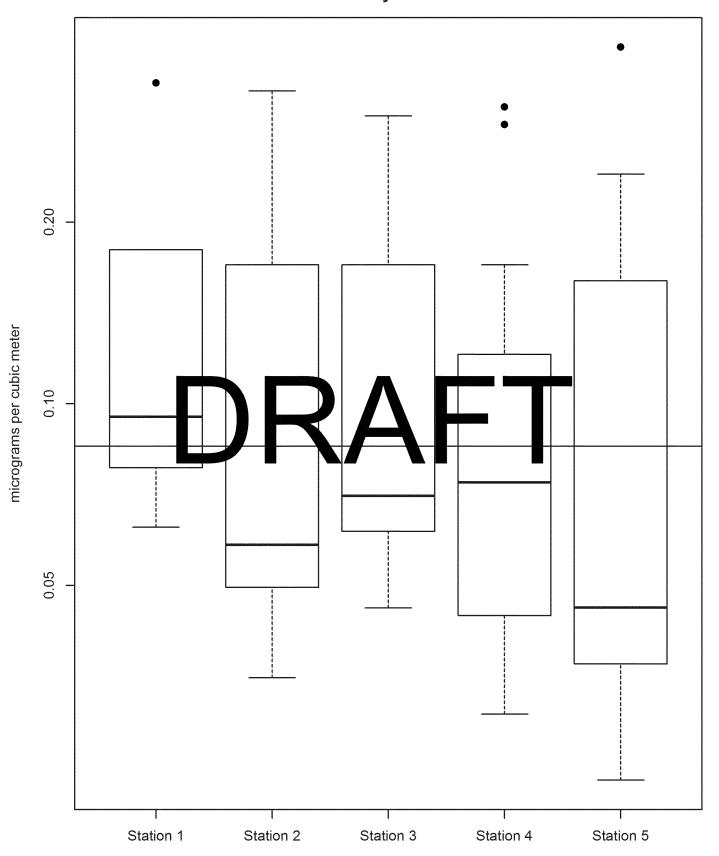


Exhibit E-5 o-Xylene





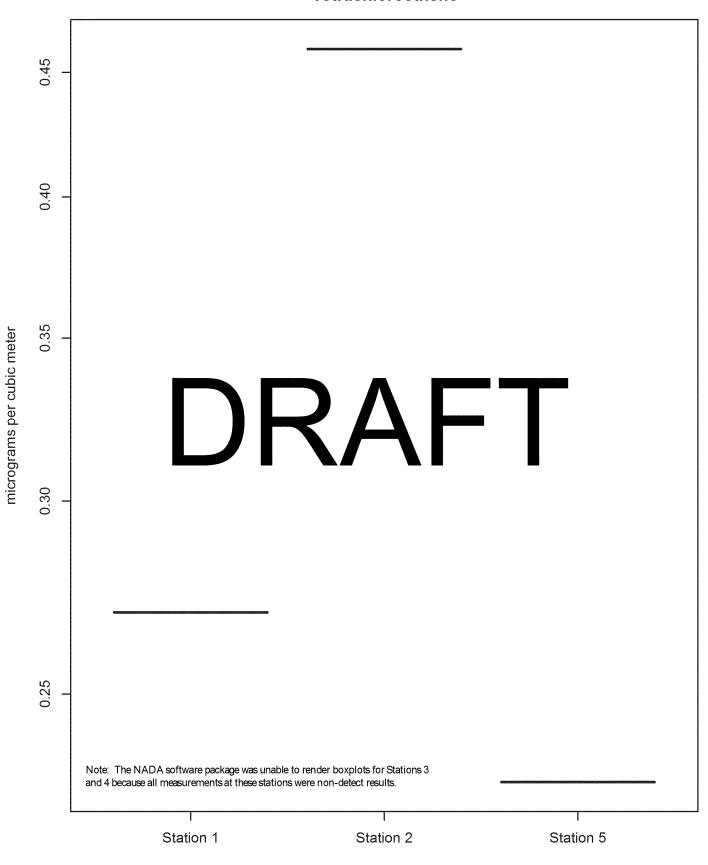


Exhibit E-7
Toluene

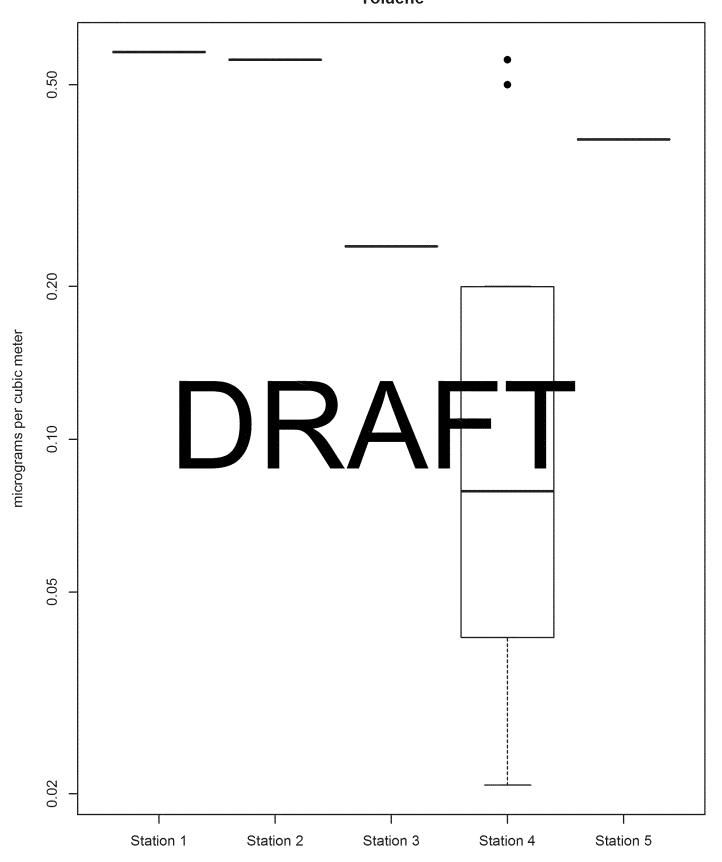
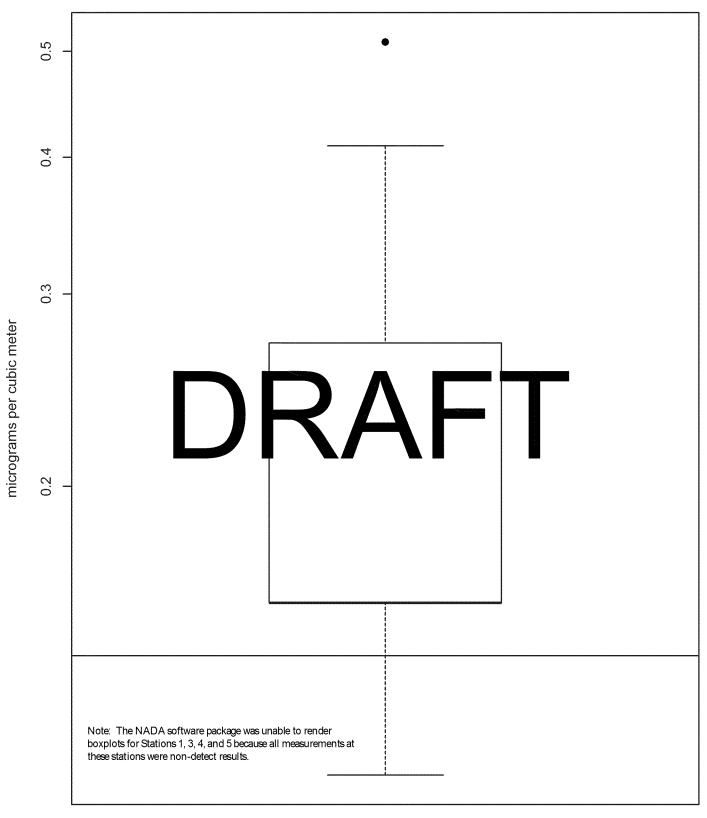


Exhibit E-8 Trichloroethene



Station 2